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(54) Title: GROUP B STREPTOCOCCUS ANTIGENS			
(57) Abstract Group B streptococcus (GBS) proteins and polynucleotides encoding them are disclosed. Said proteins are antigenic and therefore useful vaccine components for the prophylaxis or therapy of streptococcus infection in animals. Also disclosed are recombinant methods of producing the protein antigens as well as diagnostic assays for detecting streptococcus bacterial infection.			

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GROUP B STREPTOCOCCUS ANTIGENS

5

FIELD OF THE INVENTION

The present invention is related to antigens, more particularly protein antigens of group B streptococcus (GBS) 10 bacterial pathogen which are useful as vaccine components for therapy and/or prophylaxis.

BACKGROUND OF THE INVENTION

15

Streptococcus are gram (+) bacteria that are differentiated by group specific carbohydrate antigens A through O found on their cell surface. Streptococcus groups are further distinguished by type-specific capsular polysaccharide 20 antigens. Several serotypes have been identified for the Group B streptococcus (GBS) : Ia, Ib, II, III, IV, V, VI, VII and VIII. GBS also contains antigenic proteins known as "C-proteins" (alpha, beta, gamma and delta), some of which have been cloned.

25

Although GBS is a common component of the normal human vaginal and colonic flora this pathogen has long been recognized as a major cause of neonatal sepsis and meningitis, late-onset meningitis in infants, postpartum 30 endometritis as well as mastitis in dairy herds. Expectant mothers exposed to GBS are at risk of postpartum infection and may transfer the infection to their baby as the child passes through the birth canal. Although the organism is sensitive to antibiotics, the high attack rate and rapid 35 onset of sepsis in neonates and meningitis in infants results in high morbidity and mortality.

To find a vaccine that will protect individuals from GBS infection, researchers have turned to the type-specific antigens. Unfortunately these polysaccharides have proven to

5 be poorly immunogenic in humans and are restricted to the particular serotype from which the polysaccharide originates. Further, capsular polysaccharide elicit a T cell independent response i.e. no IgG production.

Consequently capsular polysaccharide antigens are unsuitable

10 as a vaccine component for protection against GBS infection.

Others have focused on the C-protein beta antigen which demonstrated immunogenic properties in mice and rabbit models. This protein was found to be unsuitable as a human

15 vaccine because of its undesirable property of interacting with high affinity and in a non-immunogenic manner with the Fc region of human IgA. The C-protein alpha antigen is rare in type III serotypes of GBS which is the serotype responsible for most GBS mediated conditions and is

20 therefore of little use as a vaccine component.

Therefore there remains an unmet need for GBS antigens that may be used as vaccine components for the prophylaxis and/or

25 therapy of GBS infection.

SUMMARY OF THE INVENTION

30 According to one aspect, the present invention provides an isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide comprising a sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,

35 SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10,

SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 15,

SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19,

SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
5 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments,
analogous or derivatives thereof.

In other aspects, there is provided vectors comprising
polynucleotides of the invention operably linked to an
10 expression control region, as well as host cells transfected
with said vectors and methods of producing polypeptides
comprising culturing said host cells under conditions
suitable for expression.

15 In yet another aspect, there is provided novel polypeptides
encoded by polynucleotides of the invention.

BRIEF DESCRIPTION OF THE DRAWINGS

20 Figure 1a is the DNA sequence of clone 1 (SEQ ID NO :1) with
corresponding amino acid sequences for open reading frames;
figure 1b is the amino acid sequence SEQ ID NO: 2;
figure 1c is the amino acid sequence SEQ ID NO: 3;
25 figure 1d is the amino acid sequence SEQ ID NO: 4;
figure 1e is the amino acid sequence SEQ ID NO: 5;
figure 1f is the amino acid sequence SEQ ID NO: 6;

Figure 2a is the DNA sequence of clone 2 (SEQ ID NO :7) with
30 corresponding amino acid sequences for open reading frames;
figure 2b is the amino acid sequence SEQ ID NO: 8;
figure 2c is the amino acid sequence SEQ ID NO: 9;
figure 2d is the amino acid sequence SEQ ID NO:10;
figure 2e is the amino acid sequence SEQ ID NO:11;
35 figure 2f is the amino acid sequence SEQ ID NO:12;

Figure 3a is the DNA sequence of clone 3 (SEQ ID NO :13) with corresponding amino acid sequences for open reading frames;

figure 3b is the amino acid sequence SEQ ID NO:14;

5 figure 3c is the amino acid sequence SEQ ID NO:15;

figure 3d is the amino acid sequence SEQ ID NO:16;

figure 3e is the amino acid sequence SEQ ID NO:17;

figure 3f is the amino acid sequence SEQ ID NO:18;

figure 3g is the amino acid sequence SEQ ID NO:19;

10 figure 3h is the amino acid sequence SEQ ID NO:20;

figure 3i is the amino acid sequence SEQ ID NO:21;

Figure 4a is the DNA sequence of clone 4 (SEQ ID NO :22) with corresponding amino acid sequences for open reading

15 frames;

figure 4b is the amino acid sequence SEQ ID NO:23;

figure 4c is the amino acid sequence SEQ ID NO:24;

figure 4d is the amino acid sequence SEQ ID NO:25;

figure 4e is the amino acid sequence SEQ ID NO:26;

20

Figure 5a is the DNA sequence of clone 5 (SEQ ID NO :27) with corresponding amino acid sequences for open reading frames;

figure 5b is the amino acid sequence SEQ ID NO:28;

25 figure 5c is the amino acid sequence SEQ ID NO:29;

figure 5d is the amino acid sequence SEQ ID NO:30;

figure 5e is the amino acid sequence SEQ ID NO:31;

Figure 6a is the DNA sequence of clone 6 (SEQ ID NO :32) ;

30 figure 6b is the amino acid sequence SEQ ID NO:33;

figure 6c is the amino acid sequence SEQ ID NO:34;

figure 6d is the amino acid sequence SEQ ID NO:35;

figure 6e is the amino acid sequence SEQ ID NO:36;

35 Figure 7a is the DNA sequence of clone 7 (SEQ ID NO :37) ;

figure 7b is the amino acid sequence SEQ ID NO:38;

figure 7c is the amino acid sequence SEQ ID NO:39;
figure 7d is the amino acid sequence SEQ ID NO:40;
figure 7e is the amino acid sequence SEQ ID NO:41;

5 Figure 8 is the DNA sequence of a part of clone 7 including
a signal sequence (SEQ ID NO :42);

Figure 9 is the DNA sequence of a part of clone 7 without a
signal sequence (SEQ ID NO :43);

10 Figure 9a is the amino acid sequence (SEQ ID NO:44);

Figure 10 represents the distribution of anti-GBS ELISA
titers in sera from CD-1 mice immunized with recombinant GBS
protein corresponding to the SEQ ID NO:39.

DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to novel antigenic polypeptides of group B streptococcus (GBS) characterized by

5 the amino acid sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,

10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,

15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments,
analogs or derivatives thereof.

A preferred embodiment of the invention includes SEQ ID NO :39 and SEQ ID NO:44.

20

A further preferred embodiment of the invention is SEQ ID NO :39.

25

A further preferred embodiment of the invention is SEQ ID NO :44.

30

As used herein, "fragments", "derivatives" or "analogs" of the polypeptides of the invention include those polypeptides in which one or more of the amino acid residues are substituted with a conserved or non-conserved amino acid residue (preferably conserved) and which may be natural or unnatural.

35

The terms «fragments», «derivatives» or «analogues» of polypeptides of the present invention also include polypeptides which are modified by addition, deletion,

substitution of amino acids provided that the polypeptides retain the capacity to induce an immune response.

By the term «conserved amino acid» is meant a substitution of one or more amino acids for another in which the antigenic determinant (including its secondary structure and hydropathic nature) of a given antigen is completely or partially conserved in spite of the substitution.

5 For example, one or more amino acid residues within the sequence can be substituted by another amino acid of a similar polarity, which acts as a functional equivalent, resulting in a silent alteration. Substitutes for an amino acid within the sequence may be selected from other members of the class to which the amino acid belongs. For example, the nonpolar (hydrophobic) amino acids include alanine, leucine, isoleucine, valine, proline, phenylalanine, tryptophan and methionine. The polar neutral amino acids include glycine, serine, threonine, cysteine, tyrosine, 10 asparagine and glutamine. The positively charged (basic) amino acids include arginine, lysine and histidine. The negatively charged (acidic) amino acids include aspartic acid and glutamic acid.

15 20 25 Preferably, derivatives and analogs of polypeptides of the invention will have about 70% identity with those sequences illustrated in the figures or fragments thereof. That is, 70% of the residues are the same. More preferably polypeptides will have greater than 95% homology. In another 30 35 preferred embodiment, derivatives and analogs of polypeptides of the invention will have fewer than about 20 amino acid residue substitutions, modifications or deletions and more preferably less than 10. Preferred substitutions are those known in the art as conserved i.e. the substituted residues share physical or chemical properties such as hydrophobicity, size, charge or functional groups.

Furthermore, in those situations where amino acid regions are found to be polymorphic, it may be desirable to vary one or more particular amino acids to more effectively mimic the 5 different epitopes of the different GBS strains.

Also included are polypeptides which have fused thereto other compounds which alter the polypeptides biological or pharmacological properties i.e. polyethylene glycol (PEG) to 10 increase half-life; leader or secretory amino acid sequences for ease of purification; prepro- and pro- sequences; and (poly)saccharides.

Moreover, the polypeptides of the present invention can be 15 modified by terminal -NH₂ acylation (eg. by acetylation, or thioglycolic acid amidation, terminal carbosy amidation, e.g. with ammonia or methylamine) to provide stability, increased hydrophobicity for linking or binding to a support or other molecule.

20 Also contemplated are hetero and homo polypeptide multimers of the polypeptide fragments, analogues and derivatives. These polymeric forms include, for example, one or more polypeptides that have been cross-linked with cross-linkers 25 such as avidin/biotin, gluteraldehyde or dimethyl-superimidate. Such polymeric forms also include polypeptides containing two or more tandem or inverted contiguous sequences, produced from multicistronic mRNAs generated by recombinant DNA technology.

30 Preferably, a fragment, analog or derivative of a polypeptide of the invention will comprise at least one antigenic region i.e. at least one epitope.

In order to achieve the formation of antigenic polymers 35 (i.e. synthetic multimers), polypeptides may be utilized having bishaloacetyl groups, nitroarylhalides, or the like,

where the reagents being specific for thio groups. Therefore, the link between two mercapto groups of the different peptides may be a single bond or may be composed of a linking group of at least two, typically at least four, 5 and not more than 16, but usually not more than about 14 carbon atoms.

In a particular embodiment, polypeptide fragments, analogs and derivatives of the invention do not contain a methionine 10 (Met) starting residue. Preferably, polypeptides will not incorporate a leader or secretory sequence (signal sequence). The signal portion of a polypeptide of the invention may be determined according to established molecular biological techniques. In general, the 15 polypeptide of interest may be isolated from a GBS culture and subsequently sequenced to determine the initial residue of the mature protein and therefor the sequence of the mature polypeptide.

20 According to another aspect, there is provided vaccine compositions comprising one or more GBS polypeptides of the invention in admixture with a pharmaceutically acceptable carrier diluent or adjuvant.

25 Suitable adjuvants include oils i.e. Freund's complete or incomplete adjuvant; salts i.e. $AlK(SO_4)_2$, $AlNa(SO_4)_2$, $AlNH_4(SO_4)_2$, $Al(OH)_3$, $AlPO_4$, silica, kaolin; saponin derivative; carbon polynucleotides i.e. poly IC and poly AU and also detoxified cholera toxin (CTB) and *E.coli* heat 30 labile toxin for induction of mucosal immunity. Preferred adjuvants include QuilATM, AlhydrogelTM and AdjuphosTM. Vaccines of the invention may be administered parenterally by injection, rapid infusion, nasopharyngeal absorption, dermoabsorption, or bucal or oral.

Vaccine compositions of the invention are used for the treatment or prophylaxis of *streptococcus* infection and/or diseases and symptoms mediated by *streptococcus* infection,

5 in particular group A *streptococcus* (*pyogenes*), group B *streptococcus* (GBS or *agalactiae*), *dysgalactiae*, *uberis*, *nocardia* as well as *Staphylococcus aureus*. General information about *Streptococcus* is available in Manual of Clinical Microbiology by P.R.Murray et al.(1995, 6th Edition, 10 ASM Press, Washington, D.C.). More particularly group B *streptococcus*, *agalactiae*. In a particular embodiment vaccines are administered to those individuals at risk of GBS infection such as pregnant women and infants for sepsis, meningitis and pneumonia as well as immunocompromised 15 individuals such as those with diabetes, liver disease or cancer. Vaccines may also have veterinary applications such as for the treatment of mastitis in cattle which is mediated by the above mentioned bacteria as well as *E.coli*.

20 The vaccine of the present invention can also be used for the manufacture of a medicament used for the treatment or prophylaxis of *streptococcus* infection and/or diseases and symptoms mediated by *streptococcus* infection, in particular group A *streptococcus* (*pyogenes*), group B *streptococcus* (GBS 25 or *agalactiae*), *dysgalactiae*, *uberis*, *nocardia* as well as *Staphylococcus aureus*. More particularly group B *streptococcus*, *agalactiae*.

Vaccine compositions are preferably in unit dosage form of 30 about 0.001 to 100 µg/kg (antigen/body weight) and more preferably 0.01 to 10 µg/kg and most preferably 0.1 to 1 µg/kg 1 to 3 times with an interval of about 1 to 12 weeks intervals between immunizations, and more preferably 1 to 6

weeks.

According to another aspect, there is provided polynucleotides encoding polypeptides of group B

5 streptococcus (GBS) characterized by the amino acid sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10,
SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15,
10 SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19,
SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24,
SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29,
SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34,
SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39,
15 SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments,
analogous or derivatives thereof.

Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a
20 (SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a
(SEQ ID NO: 37), 8 (SEQ ID NO : 42) and 9(SEQ ID NO : 43)
which correspond to the open reading frames, encoding polypeptides of the invention.

25 Preferred polynucleotides are those illustrated in figures 1a (SEQ ID NO: 1), 2a (SEQ ID NO: 7), 3a (SEQ ID NO: 13), 4a
(SEQ ID NO: 22), 5a (SEQ ID NO: 27), 6a (SEQ ID NO: 32), 7a
(SEQ ID NO: 37), 8 (SEQ ID NO : 42) and 9(SEQ ID NO : 43)
and fragments, analogues and derivatives thereof.

30 More preferred polynucleotides of the invention are those illustrated in Figures 7 (SEQ ID NO : 37), 8 (SEQ ID NO : 42) and 9(SEQ ID NO : 43).

35 Most preferred polynucleotides of the invention are those illustrated in Figures 8 (SEQ ID NO : 42) and 9(SEQ ID NO :

43).

It will be appreciated that the polynucleotide sequences illustrated in the figures may be altered with degenerate 5 codons yet still encode the polypeptides of the invention.

Due to the degeneracy of nucleotide coding sequences, other polynucleotide sequences which encode for substantially the same polypeptides of the present invention may be used in 10 the practice of the present invention. These include but are not limited to nucleotide sequences which are altered by the substitution of different codons that encode the same amino acid residue within the sequence, thus producing a silent change.

15

Accordingly the present invention further provides polynucleotides which hybridize to the polynucleotide sequences herein above described (or the complement sequences thereof) having 50% and preferably at least 70% 20 identity between sequences. More preferably polynucleotides are hybridizable under stringent conditions i.e. having at least 95% identity and most preferably more than 97% identity.

25 By capable of hybridizing under stringent conditions is meant annealing of a nucleic acid molecule to at least a region of a second nucleic acid sequence (whether as cDNA, mRNA, or genomic DNA) or to its complementary strand under standard conditions, e.g. high temperature and/or low salt 30 content, which tend to disfavor hybridization of noncomplementary nucleotide sequences. A suitable protocol is described in Maniatis T. et al., Molecular cloning : A Laboratory Manual, Cold Springs Harbor Laboratory, 1982, which is herein incorporated by reference.

35

In a further aspect, polynucleotides encoding polypeptides

of the invention, or fragments, analogs or derivatives thereof, may be used in a DNA immunization method.

That is, they can be incorporated into a vector which is replicable and expressible upon injection thereby producing

5 the antigenic polypeptide in vivo. For example polynucleotides may be incorporated into a plasmid vector under the control of the CMV promoter which is functional in eukaryotic cells. Preferably the vector is injected intramuscularly.

10

According to another aspect, there is provided a process for producing polypeptides of the invention by recombinant techniques by expressing a polynucleotide encoding said polypeptide in a host cell and recovering the expressed 15 polypeptide product. Alternatively, the polypeptides can be produced according to established synthetic chemical techniques i.e. solution phase or solid phase synthesis of oligopeptides which are ligated to produce the full polypeptide (block ligation).

20

For recombinant production, host cells are transfected with vectors which encode the polypeptide, and then cultured in a nutrient media modified as appropriate for activating promoters, selecting transformants or amplifying the genes.

25 Suitable vectors are those that are viable and replicable in the chosen host and include chromosomal, non-chromosomal and synthetic DNA sequences e.g. bacterial plasmids, phage DNA, baculovirus, yeast plasmids, vectors derived from combinations of plasmids and phage DNA. The polypeptide

30 sequence may be incorporated in the vector at the appropriate site using restriction enzymes such that it is operably linked to an expression control region comprising a promoter, ribosome binding site (consensus region or Shine-Dalgarno sequence), and optionally an operator (control element). One can select individual components of the expression control region that are appropriate for a given

host and vector according to established molecular biology principles (Sambrook et al, Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor, N.Y., 1989 incorporated herein by reference). Suitable promoters include but are not limited to LTR or SV40 promoter, *E.coli* lac, tac or trp promoters and the phage lambda P_L promoter. Vectors will preferably incorporate an origin of replication as well as selection markers i.e. ampicillin resistance gene. Suitable bacterial vectors include pET, pQE70, pQE60, pQE-9, pbs, pD10 phagescript, psIX174, pbluescript SK, pbsks, pNH8A, pNH16a, pNH18A, pNH46A, ptrc99a, pKK223-3, pKK233-3, pDR540, pRIT5 and eukaryotic vectors pBlueBacIII, pWLNEO, pSV2CAT, pOG44, pXT1, pSG, pSVK3, pBPV, pMSG and pSVL. Host cells may be bacterial i.e. *E.coli*, *Bacillus subtilis*, *Streptomyces*; fungal i.e. *Aspergillus niger*, *Aspergillus nidulans*; yeast i.e. *Saccharomyces* or eukaryotic i.e. CHO, COS.

Upon expression of the polypeptide in culture, cells are typically harvested by centrifugation then disrupted by physical or chemical means (if the expressed polypeptide is not secreted into the media) and the resulting crude extract retained to isolate the polypeptide of interest. Purification of the polypeptide from culture media or lysate may be achieved by established techniques depending on the properties of the polypeptide i.e. using ammonium sulfate or ethanol precipitation, acid extraction, anion or cation exchange chromatography, phosphocellulose chromatography, hydrophobic interaction chromatography, hydroxylapatite chromatography and lectin chromatography. Final purification may be achieved using HPLC.

The polypeptide may be expressed with or without a leader or secretion sequence. In the former case the leader may be removed using post-translational processing (see US

4,431,739; 4,425,437; and 4,338,397 incorporated herein by reference) or be chemically removed subsequent to purifying the expressed polypeptide.

5 According to a further aspect, the GBS polypeptides of the invention may be used in a diagnostic test for streptococcus infection in particular GBS infection. Several diagnostic methods are possible, for example detecting streptococcus organism in a biological sample, the following procedure may
10 be followed:

- a) obtaining a biological sample from a patient;
- b) incubating an antibody or fragment thereof reactive with a GBS polypeptide of the invention with the biological sample to form a mixture; and
- 15 c) detecting specifically bound antibody or bound fragment in the mixture which indicates the presence of streptococcus.

20 Alternatively, a method for the detection of antibody specific to a streptococcus antigen in a biological sample containing or suspected of containing said antibody may be performed as follows:

- a) isolating a biological sample from a patient;
- b) incubating one or more GBS polypeptides of the
25 invention or fragments thereof with the biological sample to form a mixture; and
- c) detecting specifically bound antigen or bound fragment in the mixture which indicates the presence of antibody specific to streptococcus.

30 One of skill in the art will recognize that this diagnostic test may take several forms, including an immunological test such as an enzyme-linked immunosorbent assay (ELISA), a radioimmunoassay or a latex agglutination assay, essentially
35 to determine whether antibodies specific for the protein are present in an organism.

The DNA sequences encoding polypeptides of the invention may also be used to design DNA probes for use in detecting the presence of streptococcus in a biological sample suspected 5 of containing such bacteria. The detection method of this invention comprises:

- a) isolating the biological sample from a patient;
- b) incubating one or more DNA probes having a DNA sequence 10 encoding a polypeptide of the invention or fragments thereof with the biological sample to form a mixture; and
- c) detecting specifically bound DNA probe in the mixture which indicates the presence of streptococcus bacteria.

15 The DNA probes of this invention may also be used for detecting circulating streptococcus i.e. GBS nucleic acids in a sample, for example using a polymerase chain reaction, as a method of diagnosing streptococcus infections. The probe may be synthesized using conventional techniques and 20 may be immobilized on a solid phase, or may be labeled with a detectable label. A preferred DNA probe for this application is an oligomer having a sequence complementary to at least about 6 contiguous nucleotides of the GBS polypeptides of the invention.

25

Another diagnostic method for the detection of streptococcus in a patient comprises:

- a) labeling an antibody reactive with a polypeptide of the invention or fragment thereof with a detectable label;
- b) administering the labeled antibody or labeled fragment 30 to the patient; and
- c) detecting specifically bound labeled antibody or labeled fragment in the patient which indicates the presence of streptococcus.

35

A further aspect of the invention is the use of the GBS

polypeptides of the invention as immunogens for the production of specific antibodies for the diagnosis and in particular the treatment of streptococcus infection.

Suitable antibodies may be determined using appropriate 5 screening methods, for example by measuring the ability of a particular antibody to passively protect against streptococcus infection in a test model. One example of an animal model is the mouse model described in the examples herein. The antibody may be a whole antibody or an antigen-10 binding fragment thereof and may in general belong to any immunoglobulin class. The antibody or fragment may be of animal origin, specifically of mammalian origin and more specifically of murine, rat or human origin. It may be a natural antibody or a fragment thereof, or if desired, a 15 recombinant antibody or antibody fragment. The term recombinant antibody or antibody fragment means antibody or antibody fragment which were produced using molecular biology techniques. The antibody or antibody fragments may be polyclonal, or preferably monoclonal. It may be specific 20 for a number of epitopes associated with the GBS polypeptides but is preferably specific for one.

EXAMPLE 1 Murine model of lethal Group B Streptococcus (GBS)
25 infection

The mouse model of GBS infection is described in detail in Lancefield et al (J Exp Med 142:165-179, 1975). GBS strain C388/90 (Clinical isolate obtained in 1990 from the 30 cephalorachidian fluid of a patient suffering from meningitis, Children's Hospital of Eastern Ontario, Ottawa, Canada) and NCS246 (National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Edmonton, Canada) were respectively serotyped as type Ia/c 35 and type II/R.

To increase their virulence, the GBS strains C388/90 (serotype Ia/c) and NCS 246 (serotype II/R) were serially passaged through mice as described previously (Lancefield et al. J Exp Med 142:165-179, 1975). Briefly, the increase of 5 virulence was monitored using intraperitoneal inoculations of serial dilutions of a subculture in Todd-Hewitt broth obtained from either the blood or spleen of infected mice. After the last passage, infected blood samples were used to inoculate Todd-Hewitt broth. After an incubation of 2 hours 10 at 37°C with 7% CO₂, glycerol at a final concentration of 10% (v/v) was added to the culture. The culture was then aliquoted and stored at -80° C for use in GBS challenge experiments. The number of cfu of GBS present in these frozen samples was determined. The bacterial concentration 15 necessary to kill 100% (LD100) of the 18 weeks old mice were determined to be 3.5X10⁵ and 1.1X10⁵ respectively for GBS strain C388/90 and NCS246, which corresponded to a significant increase in virulence for both strains. Indeed, the LD100 recorded before the passages for these two strains 20 was higher than 10⁹ cfu.

In a bacterial challenge, a freshly thawed aliquot of a virulent GBS strain was adjusted to the appropriate bacterial concentration using Todd-Hewitt broth and 1ml was 25 injected intraperitoneally to each female CD-1 mouse. The mice used for the passive protection experiments were 6 to 8 weeks old, while the ones used for the active protection experiments were approximately 18 weeks old at the time of the challenge. All inocula were verified by colony counts. 30 Animals were observed for any sign of infection four times daily for the first 48 h after challenge and then daily for the next 12 days. At the end of that period, blood samples were obtained from the survivors and frozen at -20°C. The spleen obtained from each mouse that survived the challenge 35 was cultured in order to identify any remaining GBS.

EXAMPLE 2 Immunization and protection in mice with formaldehyde killed whole GBS cells

5 Formaldehyde killed GBS whole cells were prepared according to the procedures described in Lancefield et al (J Exp Med 142:165-179, 1975). Briefly, an overnight culture on sheep blood agar plates (Quelab Laboratories, Montreal, Canada) of a GBS strain was washed twice in PBS buffer (phosphate buffered-saline, pH7.2), adjusted to approximately 3×10^9 cfu/mL and incubated overnight in PBS containing 0.3% (v/v) formaldehyde. The killed GBS suspension was washed with PBS and kept frozen at -80°C.

10

15 Female CD-1 mice, 6 to 8 weeks old (Charles River, St-Constant, Québec, Canada), were injected subcutaneously three times at two weeks interval with 0.1 ml of formaldehyde killed cells of GBS strain C388/90 ($\sim 6 \times 10^7$ GBS), or 0.1 ml of PBS for the control group. On the day before

20 the immunization, Alhydrogel™ (Superfos Biosector, Frederikssund, Denmark) at a final concentration of 0.14 mg or 0.21 mg of Al, was added to these preparations and incubated overnight at 4°C with agitation. Serum samples were obtained from each mouse before the beginning of the

25 immunization protocol and two weeks after the last injection. The sera were frozen at -20°C.

30 Eight mice in each control group injected with PBS and the group immunized with formaldehyde killed whole cells GBS strain C388/90 (Ia/c) were challenged with 1.5×10^4 cfu of GBS strain C388/90 (Ia/c) one week after the third injection. All mice immunized with the formaldehyde killed GBS whole cells survived the homologous challenge while, within 5 days after the challenge, only 4 out of the 8 mice

35 injected with PBS survived from the infection. In order to increase the mortality rate in the control groups, the

bacterial suspension had to be adjusted according to the age of the mice at the time of the bacterial challenge. In subsequent challenge experiments, when mice were older than 15 weeks, the bacterial inoculum was increased to 5 concentrations between 3.0×10^5 and 2.5×10^6 cfu.

5 Table 1 Immunization of CD1 mice with formaldehyde killed whole cells of GBS and subsequent homologous challenge [strain C388/90 (Ia/c)] and heterologous challenge [strain NCS246 (II/R)].

antigenic preparations used for immunization ¹	number of living mice 14 days after the bacterial challenge (% Survival)	
	homologous challenge: strain C388/90 (Ia/c)	heterologous challenge: strain NCS246 (II/R)
1st infection		
formaldehyde killed cells of GBS strain C388/90 (Ia/c) ²	8/8 (100) ³	n.d. ⁵
control PBS	4/8 (50)	n.d.
2nd infection		
formaldehyde killed cells of GBS strain C388/90 (Ia/c)	6/6 (100) ⁴	0/6 (0) ⁶
control PBS	2/6 (33)	0/6 (0)

¹ alhydrogel™ at a final concentration of 0.14 mg or 0.21mg of Al was used;

² approximately 6×10^7 cfu;

³ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 1.5×10^4 cfu;

⁴ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2.1×10^6 cfu;

⁵ not done;

⁶ intraperitoneal challenge with 1 mL Todd-Hewitt culture medium containing GBS NCS246 (II/R) suspension adjusted to 1.2×10^5 cfu.

In another experiment, one group of 12 mice corresponding to a control group was injected with PBS, while a second group of 12 mice was immunized with formaldehyde killed whole cells of GBS strain C388/90 (Ia/c). Six mice from each of these two groups were challenged with 2.1×10^6 cfu of the GBS strain C388/90 (Ia/c) (Table I). As the first challenge experiment, all mice immunized with the GBS strain C388/90 (Ia/c) survived the homologous challenge. Only two out of the 6 mice injected with PBS survived the infection.

The remaining 6 mice in both groups were then used one week later to verify whether this antigenic preparation could confer cross protection against strain NCS246 (II/R) 5 which produce a serologically distinct capsule. None of the mice infected with this second GBS strain survived the infection. The later result suggested that most of the protective immune response induced by formaldehyde killed strain C388/90 is directed against the capsular 10 polysaccharide and that it could be restricted to strains of that particular serotype. These results clearly indicated that this particular model of infection can be efficiently used to study the protection conferred by vaccination.

15 EXAMPLE 3 Immunization of rabbit with formaldehyde killed whole GBS cells and passive protection in mice

A New Zealand rabbit (2.5 kg, Charles River, St-Constant, 20 Québec, Canada) was immunized with formaldehyde killed cells of GBS strain C388/90 (Ia/c) to obtain hyperimmune serum. This rabbit was injected subcutaneously three times at three weeks interval with approximately 1.5×10^9 cfu of formaldehyde killed whole cells of GBS strain 25 C388/90 (Ia/c). Freund's complete adjuvant (Gibco BRL Life Technologies, Grand Island, New York) was used as the adjuvant for the first immunization, while Freund's incomplete adjuvant (Gibco BRL) was used for the following two injections. Serum samples were obtained before the 30 beginning of the immunization protocol and two weeks after the last injection. The sera were frozen at -20°C.

The ability of this particular rabbit hyperimmune serum to passively protect mice against a lethal infection with GBS

was also evaluated. Intraperitoneal injection of mice with either 15 or 25 μ L of hyperimmune rabbit serum 18 hours before the challenge protected 4 out of 5 mice (80%) against the infection. Comparatively, survival rates lower than 20% 5 were recorded for mice in the control group injected with PBS or serum obtained from a rabbit immunized with meningococcal outer membrane preparation. This result clearly indicates that the immunization of another animal species with killed GBS cells can induce the production of 10 antibodies that can passively protect mice. This reagent will also be used to characterize clones.

15 Table 2 Passive protection of CD-1 mice conferred by rabbit serum obtained after immunization with formaldehyde killed group B whole streptococci (strain C388/90 (Ia/c)) antigenic preparation

groups	number of living mice 14 days after the bacterial challenge with GBS strain C388/90 (Ia/c) ²	% survival
rabbit hyperimmune serum ² - 25 μ l	4/5	80
rabbit hyperimmune serum ¹ - 15 μ l	4/5	80
control rabbit serum - 25 μ l	1/5	20
control PBS	1/10	10

20 ¹ Freund's complete adjuvant was used for first immunization, and Freund's incomplete adjuvant for the following two injections;

² intraperitoneal challenge with 1 ml Todd-Hewitt culture medium containing GBS C388/90 (Ia/c) suspension adjusted to 2×10^4 cfu.

25

EXAMPLE 4 Recombinant production of His.Tag-GBS fusion protein

The coding region of a GBS gene was amplified by PCR (DNA 5 Thermal Cycler GeneAmp PCR system 2400 Perkin Elmer, San Jose, CA) from the genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT), respectively. The PCR product was purified from 10 agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII (Pharmacia Canada Inc Baie d'Urfe, Canada), and extracted with phenol:chloroform before ethanol precipitation. The pET-32b(+) vector (Novagen, Madison, WI) 15 containing the thioredoxin-His.Tag sequence was digested with the restriction enzymes BglII and HindIII, extracted with phenol:chloroform, and then ethanol precipitated. The BglII-HindIII genomic DNA fragment was ligated to the BglII-HindIII pET-32b(+) vector to create the coding sequence for 20 thioredoxin-His.Tag-GBS fusion protein whose gene was under control of the T7 promoter. The ligated products were transformed into *E. coli* strain XLI Blue MRF' ($\Delta(mcrA)183\Delta$ (*mcrCB-hsdSMR-mrr*)173 *endA1 supE44 thi-1 recA1 gyrA96 relA1 lac* [F' *proAB lacI^qZ Δ M15Tn10 (Tet^r)*]^c) (Stratagene, La Jolla, 25 CA) according to the method of Simanis (Hanahan, D. DNA Cloning, 1985, D.M. Glover (ed.), pp. 109-135). The recombinant pET plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing (Taq Dye Deoxy 30 Terminator Cycle Sequencing kit, ABI, Foster City, CA). The recombinant pET plasmid was transformed by electroporation (Gene Pulser II apparatus, BIO-RAD Labs, Mississauga, Canada) into *E. coli* strain AD494 (DE3) (Δ *ara-leu7697* Δ *lacX74* Δ *phoA* *PvuII* *phoR* Δ *malF3* F' [*lac^r(lacI^q) pro*] 35 *trxB::Kan* (DE3)) (Novagen, Madison, WI). In this strain of

E. coli, the T7 promoter controlling expression of the fusion protein, is specifically recognized by the T7 RNA polymerase (present on the λ DE3 prophage) whose gene is under the control of the lac promoter which is inducible by 5 isopropyl- β -D-thiogalactopyranoside (IPTG).

The transformant AD494(DE3)/rpET was grown at 37°C with agitation at 250 rpm in LB broth (peptone 10g/L, Yeast extract 5g/L, NaCl 10g/L) containing 100 μ g of ampicillin 10 (Sigma-Aldrich Canada Ltd., Oakville, Canada) per mL until the A_{600} reached a value of 0.6. In order to induce the production of the thioredoxin-His.Tag-GBS fusion protein, the cells were incubated for 2 additional hours in the presence of IPTG at a final concentration of 1mM. The 15 bacterial cells were harvested by centrifugation.

The recombinant fusion protein produced by AD494(DE3)/rpET32 upon IPTG induction for 2h was partially obtained as insoluble inclusion bodies which were purified from 20 endogenous *E. coli* proteins by the isolation of insoluble aggregates (Gerlach, G.F. et al 1992, Infect. Immun. 60:892). Induced cells from a 500 mL culture were resuspended in 20 mL of 25% sucrose-50mM Tris-HCl buffer (pH8.0) and frozen at -70°C. Lysis of cells in thawed 25 suspension was achieved by the addition of 5mL of a solution of lysozyme (10mg/mL) in 250mM Tris-HCl buffer (pH8.0) followed by an incubation of 10 to 15 min on ice, and the addition of 150mL of detergent mix (5 parts of 20mM Tris-HCl buffer [pH7.4]-300mM NaCl-2% deoxycholic acid-2% Nonidet P- 30 40 and 4 parts of 100mM Tris-HCl buffer [pH8]-50mM EDTA-2% Triton X-100) followed by 5 min incubation on ice. Upon sonication, protein aggregates were harvested by centrifugation for 30 min at 35,000 X g and a sample of the soluble cellular fraction was kept. The aggregated proteins 35 were solubilized in 6M guanidine hydrochloride. The

presence of the fusion protein in both the soluble and insoluble fractions was shown by Western Blot analysis using the serum of a mouse injected with formaldehyde killed cells of GBS strain C388/90 (Ia/c) that survived a bacterial 5 challenge with the corresponding GBS strain.

The purification of the fusion protein from the soluble fraction of IPTG-induced AD494 (DE3) /rpET was done by affinity chromatography based on the properties of the 10 His.Tag sequence (6 consecutive histidine residues) to bind to divalent cations (Ni^{2+}) immobilized on the His.Bind metal chelation resin (Novagen, Madison, WI). The purification method used are those described in the pET system Manual, 6th Edition (Novagen, Madison, WI). Briefly, the pelleted 15 cells obtained from a 100mL culture induced with IPTG was resuspended in 4mL of Binding buffer (5mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9), sonicated, and spun at 39,000 X g for 20 min to remove debris. The supernatant was filtered (0.45 μ m pore size membrane) and deposited on a column of 20 His.Bind resin equilibrated in Binding buffer. The column was then washed with 10 column volumes of Binding buffer followed by 6 column volumes of Wash buffer (20mM imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The thioredoxin-His.Tag-GBS fusion protein was eluted with Elute buffer (1M 25 imidazole-500mM NaCl-20mM Tris-HCl pH7.9). The removal of the salt and imidazole from the sample was done by dialysis against 3 X 1 liter PBS at 4°C.

30 The quantities of fusion protein obtained from either the soluble or insoluble cytoplasmic fractions of *E. coli* were estimated by Coomassie staining of a sodium dodecyl sulfate (SDS)-polyacrylamide gel with serial dilutions of these proteins and a bovine serum albumin standard (Pierce Chemical Co. Rockford, Ill.).

EXAMPLE 5 Recombinant production of GBS protein under control of lambda P_L promoter

The DNA coding region of a GBS protein was inserted 5 downstream of the promoter λP_L into the translation vector pURV22. This plasmid was derived from p629 (George et al, 1987, Bio/Technology 5:600) from which the coding region for a portion of the herpes simplex virus type I (HSV-I) glycoprotein (gD-1) was removed and the ampicillin 10 resistance gene replaced by a kanamycin cassette obtained from the plasmid vector pUC4K (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada). The vector contained a cassette of the bacteriophage λ cI857 temperature sensitive repressor gene from which the functional P_R promoter had been deleted. 15 The inactivation of the cI857 repressor by temperature increase from the ranges of 30-37°C to 37-42°C resulted in the induction of the gene under the control of λ P_L. The translation of the gene was controlled by the ribosome binding site cro followed downstream by a BglII restriction 20 site (AGATCT) and the ATG: ACTAAGGAGGTTAGATCTATG.

Restriction enzymes and T4 DNA ligase were used according to suppliers (Pharmacia Biotech Canada Inc., Baie D'Urfe, Canada; and New England Biolabs Ltd., Mississauga, Canada). 25 Agarose gel electrophoresis of DNA fragments was performed as described by Sambrook et al. (Molecular cloning : A laboratory Manual, 1989, Cold Spring Harbor Laboratory Press, N.Y). Chromosomal DNA of the GBS bacteria was prepared according to procedures described in Jayarao et al 30 (J. Clin. Microbiol., 1991, 29:2774). DNA amplification reactions by polymerase chain reaction (PCR) were made using DNA Thermal Cycler GeneAmp PCR system 2400 (Perkin Elmer, San Jose, CA). Plasmids used for DNA sequencing were purified using plasmid kits from Qiagen (Chatsworth, CA). 35 DNA fragments were purified from agarose gels using Qiaex II

gel extraction kits from Qiagen (Chatsworth, CA). Plasmid transformations were carried out by the method described by Hanahan (DNA Cloning, Glover (ed.) pp, 109-135, 1985). The sequencing of genomic DNA inserts in plasmids was done using 5 synthetic oligonucleotides which were synthesized by oligonucleotide synthesizer model 394 (the Perkin-Elmer Corp., Applied Biosystems Div. (ABI), Foster City, CA). The sequencing reactions were carried out by PCR using the Taq Dye Deoxy Terminator Cycle Sequencing kit (ABI, Foster City, CA) and DNA electrophoresis was performed on automated DNA sequencer 373A (ABI, Foster City, CA). The assembly of the 10 DNA sequence was performed using the program Sequencer 3.0 (Gene Codes Corporation, Ann Arbor, MI). Analysis of the DNA sequences and their predicted polypeptides was performed 15 with the program Gene Works version 2.45 (Intelligenetics, Inc., Mountain View CA).

The coding region of the GBS gene was amplified by PCR from GBS strain C388/90 (Ia/c) genomic DNA using oligos that 20 contained base extensions for the addition of restriction sites BglII (AGATCT) and XbaI(TCTAGA), respectively. The PCR product was purified from agarose gel using a Qiaex II gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and XbaI, and extracted with 25 phenol:chloroform before ethanol precipitation. The pURV22 vector was digested with the restriction enzymes BglII and XbaI, extracted with phenol:chloroform, and ethanol precipitated. The BglII-XbaI genomic DNA fragment was ligated to the BglII-XbaI pURV22 vector in which the GBS 30 gene was under the control of the λ PL promoter. The ligated products were transformed into *E. coli* strain XLI Blue MRF' (Δ (*mcrA*)183 Δ (*mcrCB-hsdSMR-mrr*)173 *endA1 supE44 thi-1 recA1 gyrA96 relA1 lac[F' proAB lacI^qZ Δ M15 Tn10(Tet^r)]^c) (Stratagene, La Jolla CA) according to the methods described 35 in Hanahan, supra. Transformants harboring plasmids with the*

insert were identified by analysis of lysed cells submitted to electrophoresis on agarose gel (Sambrook et al, supra). The recombinant pURV22 plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of 5 the DNA insert was verified by DNA sequencing.

The transformant XLI Blue MRF'/rpURV22 was grown at 34°C with agitation at 250 rpm in LB broth containing 50µg of kanamycin per mL until the A_{600} reached a value of 0.6. In 10 order to induce the production of the fusion protein, the cells were incubated for 4 additional hours at 39°C. The bacterial cells were harvested by centrifugation, resuspended in sample buffer, boiled for 10 min and kept at -20°C.

15

EXAMPLE 6 Subcloning GBS protein gene in CMV plasmid pCMV-GH

The DNA coding region of a GBS protein was inserted in phase 20 downstream of the human growth hormone (hGH) gene which was under the transcriptional control of the cytomegalovirus (CMV) promoter in the plasmid vector pCMV-GH (Tang et al, Nature, 1992, 356:152). The CMV promoter is non functional in E. coli cells but active upon administration of the 25 plasmid in eukaryotic cells. The vector also incorporated the ampicillin resistance gene.

The coding region of the gene was amplified by PCR from 30 genomic DNA of GBS strain C388/90 (Ia/c) using the oligos that contained base extensions for the addition of the restriction sites BglII (AGATCT) and HindIII (AAGCTT). The PCR product was purified from agarose gel using a Qiaex II 35 gel extraction kit from Qiagen (Chatsworth, CA), digested with the restriction enzymes BglII and HindIII, and extracted with phenol:chloroform before ethanol precipitation. The pCMV-GH vector (Laboratory of Dr. Stephen

A. Johnston, Department of Biochemistry, The University of Texas, Dallas, Texas) containing the human growth hormone to create fusion proteins was digested with the restriction enzymes BamHI and HindIII, extracted with phenol:chloroform, 5 and ethanol precipitated. The 1.3-kb BglII-HindIII genomic DNA fragment was ligated to the BamHI -HindIII pCMV-GH vector to create the hGH-GBS fusion protein under the control of the CMV promoter. The ligated products were transformed into *E. coli* strain DH5 α [ϕ 80 lacZ Δ M15 endA1 10 recA1 hsdR17 (r K m K s) supE44 thi-1 λ gyrA96 relA1 Δ (lacZYA-argF)U169] (Gibco BRL, Gaithersburg, MD) according to the methods described by Hanahan, supra. Transformants harboring plasmids with the insert were identified by analysis of lysed cells submitted to electrophoresis on 15 agarose gel (Sambrook, J. et al, supra). The recombinant pCMV plasmid was purified using a Qiagen kit (Qiagen, Chatsworth, CA) and the nucleotide sequence of the DNA insert was verified by DNA sequencing.

20

EXAMPLE 7 Immunological activity of GBS protein to GBS challenge

Four groups of 12 female CD-1 mice (Charles River, St- 25 Constant, Quebec, Canada) of 6 to 8 weeks were injected subcutaneously three times at three week intervals with 0.1mL of the following antigenic preparations: formaldehyde killed cells of GBS strain C388/90 (\sim 6X10 7 cfu), 20 μ g of thioredoxin-His.Tag-GBS fusion protein obtained from the 30 insoluble (inclusion bodies) or 20 μ g of the fusion protein, affinity purified (nickel column), from the soluble cytoplasmic fraction in *E.coli*, or 20 μ g of affinity purified (nickel column) thioredoxin-His.Tag control polypeptide. 20 μ g of QuillATM (Cedarlane Laboratories Ltd, Hornby, Canada)

was added to each antigenic preparation as the adjuvant. Serum samples were obtained from each mouse before immunization (PB) and on days 20 (TB1), 41 (TB2) and 54 (TB3) during the immunization protocols. Sera were frozen 5 at -20°C.

An increase of the ELISA titers was recorded after each injection of the fusion protein indicating a good primary response and a boost of the specific humoral immune response 10 after each of the second and third administration. At the end of the immunization period, the means of reciprocal ELISA titers was 456,145 for the group immunized with 20 μ g of fusion protein obtained from inclusion bodies compared to 290,133 for the group of mice immunized with the protein 15 from soluble fraction in *E.coli*. The latter result suggests that the protein obtained from inclusion bodies could be more immunogenic than the soluble protein. Analysis of mice sera in ELISA using the affinity purified thioredoxin-His.Tag to coat plates showed that negligible antibody 20 titers are made against the thioredoxin-His.Tag portion of the fusion protein. The reactivity of the sera from mice injected with the recombinant fusion protein was also tested by ELISA against formaldehyde killed whole cells of GBS strain C388/90. The antibodies induced by immunization with 25 recombinant fusion protein also recognized their specific epitopes on GBS cells indicating that their conformation is close enough to the native streptococcal protein to induce cross-reactive antibodies.

30 To verify whether the immune response induced by immunization could protect against GBS infection, mice were challenged with 3.5×10^5 cfu of GBS strains C338/90(Ia/c) and 1.2×10^5 cfu of strain NCS246(II/R) the results of which are illustrated in tables 3 and 4 respectively. Mice immunized 35 with control thioredoxin-His.Tag peptide were not protected against challenge with either GBS strain while those

immunized with formaldehyde killed C388/90 whole cells only provided protection against homologous challenge. The thioredoxin-His.Tag-GBS fusion protein of the invention protected mice from challenge with both GBS strains. Blood 5 and spleen culture of these mice did not reveal the presence of any GBS.

Table 3 Survival from GBS strain C388/90 (Ia/c) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag ²	1 / 6	17
formaldehyde killed C388/90 cells ³	6 / 6	100
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ⁴	6 / 6	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ⁴	6 / 6	100

¹ intraperitoneal administration with 1 ml Todd-Hewitt culture medium adjusted to 3.5×10^5 cfu;

² 20 μ g administered; posterior legs paralyzed in surviving mouse; GBS detected in blood and spleen;

³ 6×10^7 cfu administered;

⁴ 20 μ g administered.

Table 4 Survival from GBS strain NCS246 (II/R) challenge¹

immunizing agent	no. mice surviving challenge	% survival
thioredoxin-His.Tag ²	0 / 6	0
formaldehyde killed C388/90 cells ³	2 / 6	34
thioredoxin-His.Tag-GBS fusion (inclusion body preparation) ²	5 / 5 ⁴	100
thioredoxin-His.Tag-GBS fusion (cytoplasmic fraction) ²	6 / 6	100

5 ¹ intraperitoneal administration with 1 ml Todd-Hewitt culture medium containing GBS NCS246(II/R) suspension adjusted to 1.2×10^5 cfu.

6 ² 20 μ g administered;

7 ³ 6×10^7 cfu administered;

10 ⁴ one mouse died during immunization.

EXAMPLE 8 Immunization with recombinant GBS protein confers protection against experimental GBS infection

15

This example illustrates the protection of mice against fatal GBS infection by immunization with the recombinant protein corresponding to the SEQ ID NO:39.

20 Groups of 10 female CD-1 mice (Charles River) were immunized subcutaneously three times at three-week intervals with 20 μ g of recombinant protein purified from E. coli strain BLR (Novagen) harboring the recombinant pURV22 plasmid vector containing the GBS gene corresponding to SEQ ID NO:42 in

25 presence of 20 μ g of QuilATM adjuvant (Cedarlane Laboratories Ltd, Hornby, Canada) or, as control, with

QuilA™ adjuvant alone in PBS. Blood samples were collected from the orbital sinus on day 1, 22 and 43 prior to each immunization and fourteen days (day 57) following the third injection. One week later the mice were challenged with

5 approximately 10^4 to 10^6 CFU of various virulent GBS strains.

Samples of the GBS challenge inoculum were plated on TSA/5% sheep blood agar plates to determine the CFU and to verify the challenge dose. Deaths were recorded for a period of 14 days and on day 14 post-challenge, the surviving mice were

10 sacrificed and blood and spleen were tested for the presence of GBS organisms. The survival data are shown in table 5.

Prechallenge sera were analyzed for the presence of antibodies reactive with GBS by standard immunoassays. Elisa 15 and immunoblot analyses indicated that immunization with recombinant GBS protein produced in *E. coli* elicited antibodies reactive with both, recombinant and native GBS protein. Antibody responses to GBS are described in Example 9.

20

Table 5. Ability of recombinant GBS protein corresponding to SEQ ID NO: 39 to elicit protection against 8 diverse GBS challenge strains

5

Challenge strain				
Immunogen	Designation	Type	No. alive:	No. dead ¹
rGBS protein none	C388/90	Ia/c	8 : 2 0 : 10	(P<0.0001)
rGBS protein none	NCS 246	II/R	10 : 0 3 : 7	(P=0.0012)
rGBS protein none	ATCC12401	Ib	10 : 0 3 : 7	(P=0.001)
rGBS protein none	NCS 535	V	10 : 0 5 : 5	(P=0.01)
rGBS protein none	NCS 9842	VI	10 : 0 0 : 10	(P<0.0001)
rGBS protein NCS 915-F ³ none	NCS 915	III	7 : 3 1 : 9 4 : 6	(P=0.0007) ²
rGBS protein NCS 954-F none	NCS 954	III/R	7 : 3 4 : 6 1 : 9	(P=0.002)
rGBS protein COH1-F none	COH1	III	4 : 6 3 : 7 0 : 10	(P=0.0004)

¹ Groups of 10 mice per group were used, the number of mice surviving to infection and the number of dead mice are indicated. The survival curves corresponding to recombinant GBS protein-immunized animals were compared to the survival curves corresponding to mock-immunized animals using the log-rank test for nonparametric analysis.

² Comparison analysis to NCS915-F-immunized animals.

³ Animals were immunized with formaldehyde-killed GBS in presence of QuilA™ adjuvant.

All hemocultures from surviving mice were negative at day 14 post-challenge. Spleen cultures from surviving mice were negative except for few mice from experiment MB-11.

EXAMPLE 9 Vaccination with the recombinant GBS protein
elicits an immune response to GBS

Groups of 10 female CD-1 mice were immunized subcutaneously
5 with recombinant GBS protein corresponding to SEQ ID NO:39
as described in Example 8. In order to assess the antibody
response to native GBS protein, sera from blood samples
collected prior each immunization and fourteen days after
the third immunization were tested for antibody reactive
10 with GBS cells by ELISA using plates coated with
formaldehyde-killed GBS cells from type III strain NCS 954,
type Ib strain ATCC12401, type V strain NCS 535 or type VI
strain NCS 9842. The specificity of the raised antibodies
for GBS protein was confirmed by Western blot analyses to
15 GBS cell extracts and purified recombinant antigens. The
results shown in Figure 10 clearly demonstrate that animals
respond strongly to recombinant GBS protein used as
immunogens with median reciprocal antibody titers varying
between 12000 and 128000, for sera collected after the third
20 immunization, depending of the coating antigen. All
preimmune sera were negative when tested at a dilution of
1 :100. GBS-reactive antibodies were detectable in the sera
of each animal after a single injection of recombinant GBS
protein.

25

Example 10 Antigenic conservation of the GBS protein of the present invention

5 Monoclonal antibodies (MAbs) specific to the GBS protein of the present invention were used to demonstrate that this surface antigen is produced by all GBS and that it is also antigenically highly conserved.

10 A collection of 68 GBS isolates was used to evaluate the reactivity of the GBS-specific MAbs. These strains were obtained from the National Center for Streptococcus, Provincial Laboratory of Public Health for Northern Alberta, Canada; Centre Hospitalier Universitaire de Quebec, Pavillon CHUL, Quebec, Canada; American Type Culture Collection, USA; 15 Laboratoire de Sante Publique du Quebec, Canada; and Dept. of Infectious Disease, Children's Hospital and Medical Center, Seattle, USA. All eight MAbs were tested against the following panel of strains: 6 isolates of serotype Ia or Ia/c, 3 isolates of serotype Ib, 4 isolates of serotype II, 20 14 isolates of serotype III, 2 isolates of serotype IV, 2 isolates of serotype V, 2 isolates of serotype VI, 2 isolates of serotype VII, 1 isolate of serotype VIII, 10 isolates that were not serotyped and 3 bovine *S. agalactiae* strains. MAb 3A2 was also reacted with additional GBS: 9 isolates of serotype Ia/c and 10 isolates of serotype V. The strains were grown overnight on blood agar plates at 37°C in an atmosphere of 5% CO₂. Cultures were stored at - 25 70°C in heart infusion broth with 20% (v/v) glycerol.

30 To obtain the GBS protein-specific MAbs, mice were immunized three times at three-week intervals with 20 µg of purified recombinant GBS protein.(SEQ ID NO :44) in the presence of 20% QuillATM adjuvant. Hybridoma cell lines were generated by fusion of spleen cells recovered from immunized mice with 35 the nonsecreting SP2/0 myeloma cell line as described

previously (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Hybrid clone supernatants were tested for specific antibody production by ELISA using formaldehyde inactivated GBS and purified recombinant GBS protein (SEQ ID NO :39 or 44) as coating antigen, as previously described (Hamel, J. et al. 1987. J. Med. Microbiol. 23:163-170). Specific hybrid were cloned by limiting dilutions, expanded, and frozen in liquid nitrogen. Production of recombinant GBS protein was presented in Examples 4 & 5. Purified recombinant GBS protein or formaldehyde inactivated GBS were resolved by electrophoresis by using the discontinuous buffer system of Laemmli as recommended by the manufacturer and then transfer onto nitrocellulose membrane for Western immunoblotting as described previously (Martin et al. 1992. Infect. Immun. 60:2718-2725).

Western immunoblotting experiments clearly indicated that all eight MAbs recognized a protein band that corresponded to the purified recombinant GBS protein (SEQ ID NO :39). These MAbs also reacted with a protein band present in every GBS isolates tested so far. The reactivity of these GBS-specific MAbs are presented in Table 6. Each MAb reacted well with all 46 GBS. In addition, these MAbs also recognized the 3 *S. agalactiae* strains of bovine origin that were tested. MAb 3A2 also recognized nineteen GBS; 9 isolates of serotype Ia/c and 10 of serotype V. The other MAbs were not tested against these additional strains.

These results demonstrated that the GBS protein (SEQ ID NO :39) was produced by all the 65 GBS and the three 3 *S. agalactiae* strains of bovine origin that were tested so far. More importantly, these results clearly demonstrated that the epitopes recognized by these eight GBS-specific MAbs were widely distributed and conserved among GBS. These results also indicated that these epitopes were not

restricted to serologically related isolates since representatives of all known GBS serotypes including the major disease causing groups were tested.

- 5 In conclusion, the data presented in this example clearly demonstrated that the GBS protein of the present invention is produced by all GBS and that it is antigenically highly conserved.

10

Table 6. Reactivity of eight GBS protein-specific MAbs with different *S. agalactiae* strains as evaluated by Western immunoblots.

5

Mabs	Number of each serotype of <i>s. agalactiae</i> strains recognized by the MAbs.									NT (10) 2	TOTAL (26)	Bovine (3)
	Ia or Ia/c (6)	Ib (3)	II (4)	III (4)	IV (2)	V (2)	VI (2)	VII (2)	VIII (1)	NT (10) 2	TOTAL (26)	Bovine (3)
3A21	6	3	4	4	2	2	2	2	1	10	46	3
5A12	6	3	4	4	2	2	2	2	1	10	46	3
6G11	6	3	4	4	2	2	2	2	1	10	46	2
8B9	6	3	4	4	2	2	2	2	1	10	46	3
8E11	6	3	4	4	2	2	2	2	1	10	46	3
12B12	6	3	4	4	2	2	2	2	1	10	46	3
18F11	6	3	4	4	2	2	2	2	1	10	46	3
20G2	6	3	4	4	2	2	2	2	1	10	46	3

1 Nine additional strains of serotype Ia/c and 10 strains of serotype V were recognized by MAb 3A2.

2 These strains were not serotyped

WE CLAIM:

1. An isolated polynucleotide encoding a polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10,
SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 15,
SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19,
SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 23, SEQ ID NO: 24,
SEQ ID NO: 25, SEQ ID NO: 26, SEQ ID NO: 28, SEQ ID NO: 29,
SEQ ID NO: 30, SEQ ID NO: 31, SEQ ID NO: 33, SEQ ID NO: 34,
SEQ ID NO: 35, SEQ ID NO: 36, SEQ ID NO: 38, SEQ ID NO: 39,
SEQ ID NO: 40, SEQ ID NO: 41 and SEQ ID NO: 44 or
fragments, analogs or derivatives thereof.

2. A polynucleotide according to claim 1, wherein said polynucleotide encodes a polypeptide having at least 95% identity to the second polypeptide.

3. An isolated polynucleotide encoding a polypeptide capable of generating antibodies having binding specificity for a polypeptide having a sequence selected from the group consisting of:

SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5,
SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10,
SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 14, SEQ ID NO: 15,
SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19,
SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 23, SEQ ID NO: 24,
SEQ ID NO: 25, SEQ ID NO: 26, SEQ ID NO: 28, SEQ ID NO: 29,
SEQ ID NO: 30, SEQ ID NO: 31, SEQ ID NO: 33, SEQ ID NO: 34,
SEQ ID NO: 35, SEQ ID NO: 36, SEQ ID NO: 38, SEQ ID NO: 39,
SEQ ID NO: 40, SEQ ID NO: 41 and SEQ ID NO: 44 or
fragments, analogs or derivatives thereof.

4. An isolated polynucleotide that is complementary to the polynucleotide of claim 1.
5. An isolated polynucleotide that is complementary to the polynucleotide of claim 3.
6. The polynucleotide of claim 1, wherein said polynucleotide is DNA.
7. The polynucleotide of claim 3, wherein said polynucleotide is DNA.
8. The polynucleotide of claim 1, wherein said polynucleotide is RNA.
9. The polynucleotide of claim 3, wherein said polynucleotide is RNA.
10. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of :
SEQ ID NO : 1, SEQ ID NO : 7, SEQ ID NO : 13, SEQ ID NO : 22, SEQ ID NO : 27, SEQ ID NO : 32, SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43 or fragments, analogues or derivatives thereof.
11. A polynucleotide which hybridizes under stringent conditions to a second polynucleotide having a sequence selected from the group consisting of :
SEQ ID NO : 37, SEQ ID NO : 42 and SEQ ID NO : 43.
12. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 37.

13. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 42.
14. A polynucleotide according to claim 11 which hybridizes under stringent conditions to a second polynucleotide having the sequence SEQ ID NO : 43.
15. A polynucleotide according to claim 10 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
16. A polynucleotide according to claim 11 wherein said polynucleotide has at least 95% complementarity to the second polynucleotide.
17. A vector comprising the polynucleotide of claim 1, wherein said polynucleotide is operably linked to an expression control region.
18. A vector comprising the polynucleotide of claim 3, wherein said polynucleotide is operably linked to an expression control region.
19. A host cell transfected with the vector of claim 17.
20. A host cell transfected with the vector of claim 18.
21. A process for producing a polypeptide comprising culturing a host cell according to claim 19 under conditions suitable for expression of said polypeptide.
22. A process for producing a polypeptide comprising culturing a host cell according to claim 20 under condition suitable for expression of said polypeptide.

23. An isolated polypeptide having at least 70% identity to a second polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.
24. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 39.
25. The isolated polypeptide of claim 23 having a sequence according to SEQ ID NO : 44.
26. An isolated polypeptide capable of generating antibodies having binding specificity for a second polypeptide having a sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40, SEQ ID NO:41 and SEQ ID NO:44 or fragments, analogs or derivatives thereof.

27. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO : 39.
28. The isolated polypeptide of claim 26 having a sequence according to SEQ ID NO : 44.
29. An isolated polypeptide having an amino acid sequence selected from the group consisting of:
SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO:10, SEQ ID NO:11, SEQ ID NO:12, SEQ ID NO:14, SEQ ID NO:15, SEQ ID NO:16, SEQ ID NO:17, SEQ ID NO:18, SEQ ID NO:19, SEQ ID NO:20, SEQ ID NO:21, SEQ ID NO:23, SEQ ID NO:24, SEQ ID NO:25, SEQ ID NO:26, SEQ ID NO:28, SEQ ID NO:29, SEQ ID NO:30, SEQ ID NO:31, SEQ ID NO:33, SEQ ID NO:34, SEQ ID NO:35, SEQ ID NO:36, SEQ ID NO:38, SEQ ID NO:39, SEQ ID NO:40 and SEQ ID NO:41 or fragments, analogs or derivatives thereof.
30. The isolated polypeptide of claim 29 having an amino acid sequence according to SEQ ID NO : 39.
31. An isolated polypeptide having an amino acid sequence according to SEQ ID NO : 44.
32. An isolated polypeptide according to any one of claims 29 to 31, wherein the N-terminal Met residue is deleted.
33. An isolated polypeptide according to any one of claims 29 to 30, wherein the secretory amino acid sequence is deleted.
34. A vaccine composition comprising a polypeptide according to any one of claims 23 to 31 and a pharmaceutically acceptable carrier, diluent or adjuvant.

35. A vaccine composition comprising a polypeptide according to claim 32 and a pharmaceutically acceptable carrier, diluent or adjuvant.
36. A vaccine composition comprising a polypeptide according to claim 33 and a pharmaceutically acceptable carrier, diluent or adjuvant.
37. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 34.
38. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 35.
39. A method for therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of a composition according to claim 36.
40. A method according to any one of claims 37 to 39, wherein said animal is a bovine.
41. A method according to any one of claims 37 to 39, wherein said animal is a human.

42. A method according to any one of claims 37 to 39, wherein said bacterial infection is selected from the group consisting of group A streptococcus and group B streptococcus.
43. A method according to claim 42, wherein said bacterial infection is group B streptococcus.
44. Use of a vaccine composition according to claim 34 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
45. Use of a vaccine composition according to any one of claims 35 to 36 for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to or infected with streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
46. Use of a vaccine composition according to any one of claims 23 to 31 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.
47. Use of a vaccine composition according to claim 32 for the manufacture of a vaccine for the therapeutic or

prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

48. Use of a vaccine composition according to claim 33 for the manufacture of a vaccine for the therapeutic or prophylactic treatment of streptococcal bacterial infection in an animal susceptible to streptococcal infection comprising administering to said animal a therapeutic or prophylactic amount of the composition.

TATCTGGCAA AGAGCCAGCT AATCGTTTA GTTGGGCTAA AAATAAATTA TTAATCAATG S G K E P A N R F S W A K N K L L I N G	60
---->	
GATTCAATTGC AACTCTAGCA GCAACTATCT TATTTTTGCA AGTTCAATTG ATAGGTCTTA F I A T L A A T I L F F A V Q F I G L K	120
AACCAGATT A CCTGGAAAA ACCTACTTTA TTATCCTATT GACAGCATGG ACTTTGATGG P D Y P G K T Y F I I L L T A W T L M A	180
CATTAGTAAC TGCTTTAGTG GGATGGGATA ATAGGTATGG TTCCCTCTTG TCGTTATTAA L V T A L V G W D N R Y G S F L S L L I	240
TATTATTATT CCAGCTTGGT TCAAGCGCAG GAACTTACCC AATAGAATTG AGTCCTAAGT L L F Q L G S S A G T Y P I E L S P K F	300
TCTTCAAAC AATTCAACCA TTTTACCGA TGACTTACTC TGTTTCAGGA TTAAGAGAGA F Q T I Q P F L P M T Y S V S G L R E T	360
CCATCTCGTT GACGGGAGAC GTTAACCATC AATGGAGAAT GCTAGTAATC TTTTTAGTAT I S L T G D V N H Q W R M L V I F L V S	420
CATCGATGAT ACTTGCTCTT CTTATTTATC GTAAACAAGA AGATTAATAG AAAGTATCTA S M I L A L L I Y R K Q E D	480
GTGATAGACT AACAGTATGA TATGGTATGT CAAAGTATTT AGGAGGAGAA GATATGTCTA M S T ---->	
CTTTAACAAAT AATTATTGCA ACATTAACGT CTTTGGAAACA TTTTTATATT ATGTATTTGG L T I I I A T L T A L E H F Y I M Y L E	600
AGACGTTAGC CACCCAGTCA AATATGACTG GGAAGAGTTT TAGTATGTCT AAAGAAGAGT T L A T Q S N M T G K I F S M S K E E L	660
TGTCATATT ACCCGTTATT AAACTTTTA AGAATCAAGG TGTATACAAC GGCTTGATTG S Y L P V I K L F K N Q G V Y N G L I G	720
GCCTATTCCT CCTTTATGGG TTATATATTT CACAGAACATCA AGAAATTGTA GCTGTTTTT L F L L Y G L Y I S Q N Q E I V A V F L	780
TAATCAATGT ATTGCTAGTT GCTATTTATG GTGCTTGAC AGTTGATAAA AAAATCTTAT I N V L L V A I Y G A L T V D K K I L L	840
TAAAACAGGG TGGTTACCT ATATTAGCTC TTTAACATT CTTATTTAA TACTACTAG K Q G G L P I L A L L T F L F	900
CCGTTCGATT TAGTTGAACG GCTTTAGTA ATCATTTC TCTCATAATA CAGGTAGTTT 960	
AAGTAATTG TCTTTAAAAA TAGTATAATA TAACTACGAA TTCAAAGAGA GGTGACTTTG 1020	
ATTATGACTG AGAACTGGTT ACATACTAAA GATGGTTCAAG ATATTATTA TCGTGTGTT M T E N W L H T K D G S D I Y Y R V V ---->	
GGTCAAGGTC AACCGATTGT TTTTTACAT GGCAATAGCT TAAGTAGTCG CTATTTGAT G Q G Q P I V F L H G N S L S S R Y F D	1140
AAGCAAATAG CATATTTTC TAAGTATTAC CAAGTTATG TTATGGATAG TAGAGGGCAT K Q I A Y F S K Y Y Q V I V M D S R G H	1200
GGCAAAAGTC ATGCAAAGCT AAATACCATT AGTTTCAGGC AAATAGCAGT TGACTTAAAG G K S H A K L N T I S F R Q I A V D L K	1260

GATATCTTAG TTCATTTAGA GATTGATAAA GTTATATTGG TAGGCCATAG CGATGGTGCC 1320
 D I L V H L E I D K V I L V G H S D G A

 AATTTAGCTT TAGTTTTCA AACGATGTTT CCAGGTATGG TTAGAGGGCT TTTGCTTAAT 1380
 N L A L V F Q T M F P G M V R G L L L N

 TCAGGGAACCG TGACTATTCA TGGTCAGCGA TGGTGGGATA TTCTTTAGT AAGGATTGCC 1440
 S G N L T I H G Q R W W D I L L V R I A

 TATAAATTCC TTCACATTTT AGGGAAACTC TTTCCGTATA TGAGGCCAAA AGCTCAAGTT 1500
 Y K F L H Y L G K L F P Y M R Q K A Q V

 ATTCGCTTA TGTTGGAGGA TTTGAAGATT AGTCCAGCTG ATTTACAGCA TGTGTCAACT 1560
 I S L M L E D L K I S P A D L Q H V S T

 CCTGTAATGG TTTGGTTGG AAATAAGGAC ATAATTAAGT TAAATCATTC TAAGAAACTT 1620
 P V M V L V G N K D I I K L N H S K K L

 GCTTCTTATT TTCCAAGGGG GGAGTTTAT TCTTTAGTTG GCTTTGGCA TCACATTATT 1680
 A S Y F P R G E F Y S L V G F G H H I I

 AAGCAAGATT CCCATGTTT TAATATTATT GCAAAAAAGT TTATCAACGA TACGTTGAAA 1740
 K Q D S H V F N I I A K K F I N D T L K

 GGAGAAATTG TTGAAAAAGC TAATTGAAAA AGTCAAATCA CTGACTTCTG TGATTAAAAT 1800
 G E I V E K A N

 TGTATTTTT ATATCTGTT TAGTGCTTAT TATTGTTGAA ATGATTCAATT TGAAACGAAC 1860
 M I H L K R T
 |----->
 TATTCTGTT GAGCAACTAA AGAGTGTGTT TGGGCAATTAA TCTCCAATGA ATCTTTCTT 1920
 I S V E Q L K S V F G Q L S P M N L F L

 AATTATCCTT GTGGGGGTTA TCGCTGTCTT ACCGACAACC GGATATGACT TTGTACTGAA 1980
 I I L V G V I A V L P T T G Y D F V L N

 TGGACTTTTA CGTACAGATA AAAGCAAAAG GTATATTTA CAGACTAGTT GGTGTATCAA 2040
 G L L R T D K S K R Y I L Q T S W C I N

 CACTTTAAAT AACTTGTCAAG GATTGGTGG CTTAACGAT ATTGGGTTGC GCATGGCTTT 2100
 T F N N L S G F G G L I D I G L R M A F

 TTATGGTAAA AAAGGTCAAG AGAAGAGTGA CCTAAGAGAA GTGACTCGTT TTTTACCTA 2160
 Y G K K G Q E K S D L R E V T R F L P Y

 TCTTATTTCTT GGTCTGTCAT TTATTAGTGT GATTGCCTTA ATCATGAGCC ATATTTTCA 2220
 L I S G L S F I S V I A L I M S H I F H

 TGCCAAAGCT AGTGTGATT ACTATTATT GGTATTAATT GGTGCTAGTA TGTATTTCC 2280
 A K A S V D Y Y Y L V L I G A S M Y F P

 TGTTATTTAT TGGATTCTG GTCATAAAGG AAGCCATTAT TTCGGAGATA TGCCATCTAG 2340
 V I Y W I S G H K G S H Y F G D M P S S

 TACTCGTATA AAATTAGGTG TTGTTCTTT TTTGAATGG GGATGTGCGG CCGCAGCATT 2400
 T R I K L G V V S F F E W G C A A A A F

 TATAATTATC GGTTATTTAA TGGCATTCA TCTACCAGTT TATAAAATT TACCACTATT 2460
 I I I G Y L M G I H L P V Y K I L P L F

TTGTATTGGT TGTGCCGTCG GGATTGTATC CCTTATTCCC GGTGGATTAG GAAGTTTGA 2520
 C I G C A V G I V S L I P G G L G S F E

 ATTAGTTCTA TTTACAGGGT TTGCTGCCGA GGGACTACCT AAAGAAACTG TGGTTGCATG 2580
 L V L F T G F A A E G L P K E T V V A W

 GTTATTACTT TATCGTTAG CCTACTATAT TATTCCATTC TTTGCAGGTA TCTATTCTT 2640
 L L L Y R L A Y Y I I P F F A G I Y F F

 TATCCATTAT TTAGGTAGTC AAATAAATCA ACGBTATGAA AATGTCGGAA AAGAGTTAGT 2700
 I H Y L G S Q I N Q R Y E N V P K E L V

 ATCAACTGTT CTACAAACCA TGGTGAGCCA TTTGATGCGT ATTTAGGTG CATTCTTAAT 2760
 S T V L Q T M V S H L M R I L G A F L I
 |---->
 ATTTCAACA GCATTTTTG AAAATATTAC TTATATTATG TGGTTGCAGA AGCTAGGCTT 2820
 F S T A F F E N I T Y I M W L Q K L G L

 GGACCCATTA CAAGAACAAA TGTATGGCA GTTCCAGGT TTATTGCTGG GGGTTTGT 2880
 D P L Q E Q M L W Q F P G L L L G V C F

 TATTCTCTTA GCTAGAACTA TTGATCAAAA AGTGAATGAA GCTTTCCAA TTGCTATTAT 2940
 I L L A R T I D Q K V K N A F P I A I I

 CTGGATTACT TTGACATTGT TTTATCTTAA TTTAGGTCA ATTAGTTGGC GACTATCTT 3000
 W I T L T L F Y L N L G H I S W R L S F

 CTGGTTTATT TTACTATTGT TAGGCTTATT AGTCATTAAG CCAACTCTCT ATAAAAAAACA 3060
 W F I L L L G L L V I K P T L Y K K Q

 ATTTATTTAT AGCTGGGAAG AGCGTATTAA GGATGGAATC ATTATCGTT GTTAATGGG 3120
 F I Y S W E E R I K D G I I I V S L M G

 AGTTCTATT TATATTGCAG GACTACTATT CCCTATCAGG GCTCATATTA CAGGTGGTAG 3180
 V L F Y I A G L L F P I R A H I T G G S

 TATTGAACGC CTGCATTATA TCATAGCATG GGAGCCGATA GCATTGGCTA CGTTGATTCT 3240
 I E R L H Y I I A W E P I A L A T L I L

 TACTCTCGTT TATTTATGTT TGGTTAAGAT TTTACAAGGA AAATCTTGTG AGATTGGTGA 3300
 T L V Y L C L V K I L Q G K S C Q I G D

 TGTGTTCAAT GTGGATCGTT ATAAAAAAACT ACTTCAAGCT TACGGTGGTT CTTGGATAG 3360
 V F N V D R Y K K L L Q A Y G G S S D S

 CGGTTTAGCC TTTTAAATG ATAAAAGGCT CTACTGGTAC CAAAAAAATG GAGAAGATTG 3420
 G L A F L N D K R L Y W Y Q K N G E D C

 CGTTGCGTTC CAATTTGTA TTGTCATAA TAAATGTCTT ATTATGGGG AACCCAGCCGG 3480
 V A F Q F V I V N N K C L I M G E P A G

 TGATGACACT TATATTGCGT AAGCTATTGA ATCGTTATT GATGATGCTG ATAAGCTAGA 3540
 D D T Y I R E A I E S F I D D A D K L D

 CTATGACCTT GTTTTACCA GTATTGGACA GAAGTTGACA CTACTTTAC ATGAGTATGG 3600
 Y D L V F Y S I G Q K L T L L L H E Y G

 TTTGACTTT ATGAAAGTTG GTGAGGATGC TTTAGTTAAT TTAGAAACGT TTACTCTTAA 3660
 F D F M K V G E D A L V N L E T F T L K

AGGGATAAG TACAAACCTT TCAGAAATGC CCTAAATAGA GTTGAAAAGG ATGGTTCTA	3720
G N K Y K P F R N A L N R V E K D G F Y	
TTTCGAAGTT GTACAATCGC CACATAGTCA AGAGCTACTA AATAGTTGG AAGAGATTC	3780
F E V V Q S P H S Q E L L N S L E E I S	
TAATACTTGG TTAGAAGGAC GTCTGAAAA AGGTTCTCA CTAGGATATT TTAATAAAGA	3840
N T W L E G R P E K G F S L G Y F N K D	
TTATTTCCAA CAAGCCCCAA TAGCTTGGT AAAAAATGCT GAACACGAAG TTGTTGCTTT	3900
Y F Q Q A P I A L V K N A E H E V V A F	
TGCTAATATT ATGCCAAACT ATGAAAAGAG TATTATCTCT ATTGATTAA TGGTCACGA	3960
A N I M P N Y E K S I I S I D L M R H D	
TAAACAGAAA ATTCCGAATG GCGTTATGGA TTTCTCTTT TTATCATTAT TCTCTTATTA	4020
K Q K I P N G V M D F L F L S L F S Y Y	
TCAAGAGAAAG GGATACCACT ATTTGATTT GGGGATGGCA CCTTTATCAG GAGTTGGTCG	4080
Q E K G Y H Y F D L G M A P L S G V G R	
CGTTGAAACA AGTTTGCTA AAGAGAGAAT GGCATATCTT GTCTATCATT TCGGTAGTCA	4140
V E T S F A K E R M A Y L V Y H F G S H	
TTTCTACTCA TTTAATGGTT TACACAAGTA TAAGAAGAAG TTTACACCAT TGTGGTCGGA	4200
F Y S F N G L H K Y K K K F T P L W S E	
ACGTTATATT TCTTGTCTC GTTGTCTG GTTAATTGT GCTATTGTG CCCTATTAAT	4260
R Y I S C S R S S W L I C A I C A L L M	
GGAAGATAGT AAAATTAAGA TTGTTAAATA AGCTTTATTT GGCAATTAAA AAGAGCATGT	4320
E D S K I K I V K	
CATGCGACAT GCTTTTTA AATCATTTAA TACCATGAT TGCTGAATC TACTTATAA	4380
TATGATGTGC TTTAAATAT TGTTAGCTA CTGTAGCTGC TGATTATGC TTTACAGCTA	4440
CTTGGTAGTT CATTCTTGC ATTCTTTT CAGTGATATG ACCAGCAAGT TTATTGAGAG	4500
CTTTTTTAC TTGA (SEQ ID NO:1)	4514

FIG. 1a
[clone1-dna/aa]

SGKEPANRFS WAKNKLLING FIATLAATIL FFAVQFIGLK PDYPGKTYFI 50
ILLTAWTLMA LVTALVGWDN RYGSFLSLLI LLFQLGSSAG TYPIELSPKE 100
FQTIQPFPLPM TYSVSGLRET ISLTGDVNHQ WRMLVIFLVS SMILALLIYR 150
KQED (SEQ ID NO:2) 154

FIG. 1b

MSTLTIIIAAT LTALEHFYIM YLETLATQSN MTGKIFSMSK EELSYLPVIK 50
LFKNQGVYNG LIGLFLLYGL YISQNQEIVA VFLINVLLVA IYGALTVDKK 100
ILLKQGGLPI LALLTFLF (SEQ ID NO:3) 118

FIG. 1c

MTENWLHTKD GSDIYYRVVG QGQPIVFLHG NSLSSRYFDK QIAYFSKYYQ 50
VIVMDSRGHG KSHAKLNTIS FRQIAVDLKD ILVHLEIDKV ILVGHSDGAN 100
LALVFQTMFP GMVRGLLLNS GNLTIHQRW WDILLVRIAY KFLHYLGKLF 150
PYMRQKAQVI SLMLEDLKIS PADLQHVSTP VMVLVGNKDI IKLNHSKKLA 200
SYFPRGEFYS LVGF GHIIK QDSHVFNIIA KKFINDTLKG EIVEKAN 247
(SEQ ID NO:4)

FIG. 1d

MIHLKRTISV EQLKSVFGQL SPMNLFILIL VGVIAVLPTT GYDFVLNGLL	50
RTDKSKRYIL QTSWCINTFN NLSGFGGLID IGLRMAFYK KGQEKSRLRE	100
VTRFLPYLIS GLSFISVIAL IMSHIFHAKA SVDYYLVLI GASMYFPVIY	150
WISGHKGSHY FGDMPSSTRI KLGVVSSFEW GCAAAAFIII GYLMGIHLPV	200
YKILPLFCIG CAVGIVSLIP GGLGSFELVL FTGFAAEGLP KETVVAWLLL	250
YRLAYYIIPF FAGIYFFIHY LGSQINQRYE NVPKELVSTV LQTMVSHLMR	300
ILGAFLIFST AFFENITYIM WLQKLGDLPL QEQMLWQFPG LLLGVCFILL	350
ARTIDQKVKN AFPIAIIWIT LTLFYLNLGH ISWRLSFWFI LLLLGLLVIK	400
PTLYKKQFIY SWEERIKDGI IIVSLMGVLF YIAGLLFPIR AHITGGSIER	450
LHYIIIAWEPI ALATLILTLV YLCLVKILQG KSCQIGDVFN VDRYKKLLQA	500
YGGSSDGLA FLNDKRLYWY QKNGEDCVAF QFVIVNNKCL IMGEPAGDDT	550
YIREAIESFI DDADKLDYDL VFYSIGQKLT LLLHEYGFDF MKVGEDALVN	600
LETFTLKGK YKPFERNALNR VEKDGFYFEV VQSPHSQELL NSLEEISNTW	650
LEGRPEKGFS LGYFNKDYFQ QAPIALVKNA EHEVVAFANI MPNYEKSIIS	700
IDLMRHDKQK IPNGVMDLF LSLFSYYQEK GYHYFDLGMA PLSGVGRVET	750
SFAKERMAYL VYHFGSHFYS FNGLHKYKKK FTPLWSERYI SCSRSSLIC	800
AICALLMEDS KIKIVK (SEQ ID NO:5)	816

FIG. 1e

MRILGAFLIF STAFFENITY IMWLQKLGKD PLQEQMLWQF PGLLLGVCFI	50
LLARTIDQKV KNAFPIAIW ITLTLFYLNL GHISWRLSFW FILLLLGLLV	100
IKPTLYKKQF IYSWEERIKD GIIIVSLMGV LFYIAGLLFP IRAHITGGSI	150
ERLHYIIIAWE PIALATLILT LVYLCLVKIL QGKSCQIGDV FNVDRYKKLL	200
QAYGGSSDGLA LAFLNDKRLY WYQKNGEDCV AFQFVIVNNK CLIMGEPAGD	250
DTYIREAIES FIDDADKLDY DLVFYSIGQK LTLLLHEYGF DFMKGEDAL	300
VNLETFTLKG NKYPFRNAL NRVEKDGFYF EVVQSPHSQE LLNSLEEISN	350
TWLEGRPEKG FSLGYFNKDY FQQAPIALVK NAEHEVVAFA NIMPNEYKSI	400
ISIDLMRHDK QKIPNGVMDF LFLSLFSYYQ EKGYHYFDLG MAPLSGVGRV	450
ETSFAKERMA YLVYHFGSHF YSFNGLHKYK KKFTPLWSER YISCSRSSL	500
ICAICALLME DSKIKIVK (SEQ ID NO:6)	518

FIG. 1f

AATTTTGATA	TCGAAACAAC	AACTTTGAG	GCAATGAAAA	AGCACCGC	TC ATTATTGGAG	60
N F D I	E T T	T F E	A M K K	H A S	L L E	
---->						
AAAATATCTG	TTGAGCGTTC	TTTTATTGAA	TTTGATAAAC	TTCTATTAGC	ACCTTATTGG	120
K I S V	E R S	F I E	F D K L	L L A	P Y W	
CGTAAAGGAA	TGCTGGCACT	AATAGATAGT	CATGCTTTA	ATTATCTACC	ATGCTTAAAA	180
R K G M	L A L	I D S	H A F N	Y L P	C L K	
AATAGGGAAT	TACAATTAAG	CGCCCTTTTG	TCCCAGTTAG	ATAAAGATT	TTTATTGAG	240
N R E L	Q L S	A F L	S Q L D	K D F	L F E	
ACATCAGAAC	AAGCTGGGC	ATCACTCATC	TTGAGTATGG	AAGTTGAACA	CACAAAGACT	300
T S E Q	A W A	S L I	L S M E	V E H	T K T	
TTTTAAAAAA	AATGGAAGAC	ATCAACTCAC	TTTCAAAAAG	ATGTTGAGCA	TATAGTGGAT	360
F L K K	W K T	S T H	F Q K D	V E H	I V D	
GT T T A T C G T A	TTCGTGAACA	AATGGGATTG	GCTAAAGAAC	ATCTTTATCG	TTATGGAAAA	420
V Y R I	R E Q	M G L	A K E H	L Y R	Y G K	
ACTATAATAA	AACAAGCGGA	AGGTATT CGC	AAAGCAAGAG	GCTTGATGGT	TGATTCGAA	480
T I I K	Q A E	G I R	K A R G	L M V	D F E	
AAAATAGAAC	AACTAGATAG	TGAGTTAGCA	ATCCATGATA	GGCATGAGAT	AGTTGTCAAT	540
K I E Q	L D S	E L A	I H D R	H E I	V V N	
GGTGGCACCT	TAATCAAGAA	ATTAGGAATA	AAACCTGGTC	CACAGATGGG	AGATATTATC	600
G G T L	I K K	L G I	K P G P	Q M G	D I I	
TCTCAAATTG	AATTAGCCAT	TGTTTTAGGA	CAACTGATTA	ATGAAGAAGA	GGCTATTTA	660
S Q I E	L A I	V L G	Q L I N	E E E	A I L	
CATTTGTTA	AGCAGTACTT	GATGGATTAG	AGAGGATTAT	ATGAGCGATT	TTTTAGTAGA	720
H F V K	Q Y L	M D		M S D F	L V D	
---->						
TGGATTGACT	AAGTCGGTTG	GTGATAAGAC	GGTCTTTAGT	AATGTTCAT	TTATCATCCA	780
G L T	K S V G	D K T	V F S	N V S F	I I H	
TAGTTTAGAC	CGTATTGGGA	TTATTGGTGT	CAATGGAAC	GGAAAGACAA	CACTATTAGA	840
S L D	R I G I	I G V	N G T	G K T T	L L D	
TGTTATTCG	GGTGAATTAG	GT T T G A T G G	TGATCGTTCC	CCTTTTCAT	CAGCTAATGA	900
V I S	G E L G	F D G	D R S	P F S S	A N D	
TTATAAGATT	GCTTATTTAA	AACAAGAAC	AGACTTTGAT	GATTCTCAGA	CAATTTGGA	960
Y K I	A Y L K	Q E P	D F D	D S Q T	I L D	
CACCGTACTT	TCTTCTGACT	TAAGAGAGAT	GGCTTTAATT	AAAGAATATG	AATTATTGCT	1020
T V L	S S D L	R E M	A L I K	E Y E	L L L	
TAATCACTAC	GAAGAAAGTA	ACCAATCACG	TCTAGAGAAA	GTAATGGCAG	AAATGGATT	1080
N H Y	E E S K	Q S R	L E K	V M A E	M D S	
TTTAGATGCT	TGGTCTATTG	AGAGCGAAGT	CAAAACAGTA	TTATCCAAT	TAGGTATTAC	1140
L D A	W S I E	S E V	K T V	L S K L	G I T	
TGATTTGCAG	TTGTCGGTTG	GTGAATTATC	AGGAGGATTA	CGAAGACGTG	TTCAATTAGC	1200
D L Q	L S V G	E L S	G G L	R R R V	Q L A	

GCAAGTATTAA TTAAATGATG CAGATTTATT GCTCTTAGAC GAACCTACTA ACCACTTAGA 1260
 Q V L L N D A D L L L L D E P T N H L D

 TATTGACACT ATTGCATGGT TAACGAATT TTTGAAAAAT AGTAAAAGA CAGTGCTTT 1320
 I D T I A W L T N F L K N S K K T V L F

 TATAACTCAT GATCGTTATT TTCTAGACAA TGTTGCAACA CGTATTTTG AATTAGATAA 1380
 I T H D R Y F L D N V A T R I F E L D K

 GGCACAGATT ACAGAATATC AAGGCAATT TCAGGATTAT GTCCGACTTC GTGCAGAAC 1440
 A Q I T E Y Q G N Y Q D Y V R L R A E Q

 AGACGAGCGT GATGCTGCTA GTTTACATAA AAAGAAACAG CTTTATAAAC AGGAACCTAGC 1500
 D E R D A A S L H K K K Q L Y K Q E L A

 TTGGATGCGT ACTCAGCCAC AAGCTCGTGC AACGAAACAA CAGGCTCGTA TTAATCGTT 1560
 W M R T Q P Q A R A T K Q Q A R I N R F

 TCAAAATCTA AAAAACGATT TACACCAAC AAGCGATACA AGCGATTTGG AAATGACATT 1620
 Q N L K N D L H Q T S D T S D L E M T F

 TGAAACAAAGT CGAATTGGGA AAAAGGTTAT TAATTTGAA AATGTCCTT TTTCTTACCC 1680
 E T S R I G K K V I N F E N V S F S Y P

 AGATAAAATCT ATCTTGAAAG ACTTTAATT GTTAATTCAA AATAAAGACC GTATTGGCAT 1740
 D K S I L K D F N L L I Q N K D R I G I

 CGTTGGAGAT AATGGTGTG GAAAGTCAAC CTTACTTAAT TTAATTGTT AAGATTTACA 1800
 V G D N G V G K S T L L N L I V Q D L Q

 GCCGGATTCTG GGTAATGTCT CTATTGGTGA AACGATACGT GTAGGTTACT TTTCACAACA 1860
 P D S G N V S I G E T I R V G Y F S Q Q

 ACTTCATAAT ATGGATGGCT CAAAACGTGT TATTAATTAT TTGCAAGAGG TTGCAGATGA 1920
 L H N M D G S K R V I N Y L Q E V A D E

 GGTTAAAACT AGTGTGGTA CAACAAGTGT GACAGAACTA TTGGAACAAT TTCTCTTCC 1980
 V K T S V G T T S V T E L L E Q F L F P

 ACGTTCGACA CATGGAACAC AAATTGCAAA ATTATCAGGT GGTGAGAAAA AAAGACTTTA 2040
 R S T H G T Q I A K L S G G E K K R L Y

 CCTTTAAAAA ATCCTGATTG AAAAGCTAA TGTGTTACTA CTTGATGAGC CGACAAATGA 2100
 L L K I L I E K P N V L L L D E P T N D

 CTTAGATATT GCTACATTAA CTGTTCTGAA AAATTTTTA CAAGGCTTGC GTGGTCCTGT 2160
 L D I A T L T V L E N F L Q G F G G P V

 GATTACAGTT AGTCACGATC GTTACTTTT AGATAAAGTG GCTAATAAAA TTATTGCGTT 2220
 I T V S H D R Y F L D K V A N K I I A F

 TGAAGATAAC GATATCCGTG AATTTTTGG TAATTATACT GATTATTTAG ATGAAAAAGC 2280
 E D N D I R E F F G N Y T D Y L D E K A

 ATTTAATGAG CAAAATAATG AAGTTATCAG TAAAAAAGAG AGTACCAAGA CAAGTCGTGA 2340
 F N E Q N N E V I S K K E S T K T S R E

 AAAGCAAAGT CGTAAAAGAA TGTCTTACTT TGAAAAACAA GAATGGCGA CAATTGAAGA 2400
 K Q S R K R M S Y F E K Q E W A T I E D

 CGATATTATG ATATTGGAAA ATACTATCAC TCGTATAGAA AATGATATGC AAACATGTGG 2460

D I M I L E N T I T R I E N D M Q T C G	
TAGTGATTT ACAAGGTTAT CTGATTTACA AAAGGAATTA GATGCAAAAAA ATGAAGCACT	2520
S D F T R L S D L Q K E L D A K N E A L	
TCTAGAAAAG TATGACCGTT ATGAGTACCT TAGTGAGTTA GACACATGAT TATCCGTCCG	2580
L E K Y D R Y E Y L S E L D T M I I R P	
ATTATTAAGG ATGATGACCA AGCAGTTGCA CAATTAATTG GACAAAGTTT ACGCGCCTAT	2640
I I K N D D Q A V A Q L I R Q S L R A Y	
GATTTAGATA AACCTGATAC AGCATATTCA GACCCTCACT TAGATCATT GACCTCATAAC	2700
D L D K P D T A Y S D P H L D H L T S Y	
TACGAAAAAA TAGAGAAGTC AGGATTCTTT GTCATTGAGG AGAGAGATGA GATTATTGGC	2760
Y E K I E K S G F F V I E E R D E I I G	
TGTGGCGGCT TTGGTCCGCT GAAAAATCTA ATTGCAGAGA TGCAGAAGGT GTACATTGCA	2820
C G G F G P L K N L I A E M Q K V Y I A	
GAACGTTTCC GTGGTAAGGG GCTTGCTACT GATTTAGTGA AAATGATTGA AGTAGAAGCT	2880
E R F R G K G L A T D L V K M I E V E A	
CGAAAAATTG GGTATAGACA ACTTTATTTA GAGACAGCCA GTACTTTGAG TAGGGCAACT	2940
R K I G Y R Q L Y L E T A S T L S R A T	
GCGGTTTATA AGCATATGGG ATATTGTGCC TTATCGCAAC CAATAGCAAA TGATCAAGGT	3000
A V Y K H M G Y C A L S Q P I A N D Q G	
CATACAGCTA TGGATATTTG GATGATTAAA GATTTATAAG TTGAAAGTGG ATTAGTGAAC	3060
H T A M D I W M I K D L	
ATGGATTAAT TATTTGAGA TAAGAGGAAA GAAAAGGAGA CATATATGGC ATATATTGG	3120
M A Y I W	
TCTTATTTGA AAAGGTACCC CAATTGGTTA TGGCTTGATT TACTAGGAGC TATGCTTTTT	3180
S Y L K R Y P N W L W L D L L G A M L F	
GTGACGGTTA TCCTAGGAAT GCCCACAGCC TTAGCGGGTA TGATTGATAA TGCGCTTACA	3240
V T V I L G M P T A L A G M I D N G V T	
AAAGGTGATC GGACTGGAGT TTATCTGTGG ACGTTCATCA TGTTTATATT TGGTGTACTA	3300
K G D R T G V Y L W T F I M F I F V V L	
GGTATTATTG GGC GTATTAC GATGGCTTAC GCATCTAGTC GCTTAACGAC AACAAATGATT	3360
G I I G R I T M A Y A S S R L T T T M I	
AGAGATATGC GTAATGATAT GTATGCTAAG CTTCAAGAAT ACTCCCATCA TGAATATGAA	3420
R D M R N D M Y A K L Q E Y S H H E Y E	
CAGATAGGTG TATCTTCACT AGTGACACGT ATGACAAGCG ATACTTTGT TTTGATGCAA	3480
Q I G V S S L V T R M T S D T F V L M Q	
TTTGCTGAAA TGTCTTACG TTTAGGCCTA GTAACTCCTA TGGTAATGAT TTTAGCGTG	3540
F A E M S L R L G L V T P M V M I F S V	
GTTATGATAC TAATTACGAG TCCATCTTG GCTTGGCTTG TAGCGGTTGC GATGCCTCTT	3600
V M I L I T S P S L A W L V A V A M P L	
TTGGTAGGAG TCGTTTATA TGTAGCTATA AAAACAAAAC CTTTATCTGA AAGACAACAG	3660
L V G V V L Y V A I K T K P L S E R Q Q	

ACTATGTTG ATAAAATCAA TCAATATGTT CGTAAAAATT TAACAGGGTT ACGCGTTGTT	3720
T M L D K I N Q Y V R E N L T G L R V V	
AGAGCCTTG CAAGAGAGAA TTTCAATCA CAAAAATTTC AAGTCGCTAA CCAACGTTAC	3780
R A F A R E N F Q S Q K F Q V A N Q R Y	
ACAGATACTT CAACTGGTCT TTTAAATTA ACAGGGCTAA CAGAACCACT TTTCGTTCAA	3840
T D T S T G L F K L T G L T E P L F V Q	
ATTATTATTG CAATGATTGT GGCTATCGTT TGGTTTGCTT TGGATCCCTT ACAAAAGAGGT	3900
I I I A M I V A I V W F A L D P L Q R G	
GCTATTAAAA TAGGGGATTT AGTGCTTT ATCGAATATA GCTTCATGC TCTCTTTCA	3960
A I K I G D L V A F I E Y S F H A L F S	
TTTTGCTAT TTGCCAATCT TTTACTATG TATCCTCGTA TGGTGGTATC AAGCCATCGT	4020
F L L F A N L F T M Y P R M V V S S H R	
ATTAGAGAGG TGATGGATAT GCCAATCTCT ATCAATCTTA ATGCCGAAGG TGTTACGGAT	4080
I R E V M D M P I S I N P N A E G V T D	
ACGAAACTTA AAGGGCATT AGAATTGAT AATGTAACAT TCGTTATCC AGGAGAAACA	4140
T K L K G H L E F D N V T F A Y P G E T	
GAGAGTCCCG TTTGCATGA TATTCTTTT AAAGCTAACG CTGGAGAAC AATTGCTTTT	4200
E S P V L H D I S F K A K P G E T I A F	
ATTGGTTCAA CAGGTTCAAG AAAATCTCT CTTGTTAATT TGATTCCACG TTTTATGAT	4260
I G S T G S G K S S L V N L I P R F Y D	
GTGACACTTG GAAAATCTT AGTAGATGGA GTTGATGTA GAGATTATAA CCTTAAATCA	4320
V T L G K I L V D G V D V R D Y N L K S	
CTTCGCCAAA AGATTGGATT TATCCCCAA AAAGCTCTT TATTACAGG GACAATAGGA	4380
L R Q K I G F I P Q K A L L F T G T I G	
GAGATTAA AATATGGAAA AGCTGATGCT ACTATTGATG ATCTTAGACA AGCGGTTGAT	4440
E N L K Y G K A D A T I D D L R Q A V D	
ATTTCTCAAG CTAAAGAGTT TATTGAGAGT CACCAAGAAG CCTTGAAAC GCATTTAGCT	4500
I S Q A K E F I E S H Q E A F E T H L A	
GAAGGTGGGA GCAATCTTC TGGGGTCAA AAACAACGGT TATCTATTGC TAGGGCTGTT	4560
E G G S N L S G G Q K Q R L S I A R A V	
GTAAAGATC CAGATTATA TATTTTGAT GATTCATTCT CGCTCTCGA TTATAAGACA	4620
V K D P D L Y I F D D S F S A L D Y K T	
GACGCTACTT TAAGAGCGCG TCTAAAAGAA GTAACCGGT ATTCTACAGT TTTGATAGTT	4680
D A T L R A R L K E V T G D S T V L I V	
GCTCAAAGGG TGGGTACGAT TATGGATGCT GATCAGATTA TTGTCTTGA TGAAGGGCAA	4740
A Q R V G T I M D A D Q I I V L D E G E	
ATTGTCGGTC GTGGTACCCA CGCTCAATTA ATAGAAAATA ATGCTATTAA TCGTGAAATC	4800
I V G R G T H A Q L I E N N A I Y R E I	
GCTGAGTCAC AACTGAAGAA CCAAAACTTA TCAGAAGGAG AGTGATTGTA TGAGAAAAAA	4860
A E S Q L K N Q N L S E G E M R K K	

|---->

ATCTGTTTT TTGAGATTAT GGTCTTACCT AACTCGCTAC AAAGCTACTC TTTTCTTAGC	4920
S V F L R L W S Y L T R Y K A T L F L A	
GATTTTTTG AAAGTTTAT CTAGTTTAT GAGTGTCTG GAGCCTTTA TTTTAGGGTT	4980
I F L K V L S S F M S V L E P F I L G L	
AGCGATAACA GAGTTGACTG CTAACCTTGT TGATATGGCT AAGGGAGTTT CTGGGGCAGA	5040
A I T E L T A N L V D M A K G V S G A E	
ATTGAACGTT CCTTATATTG CTGGTATTG GATTATTAT TTTTCAGAG GTGTTTCTA	5100
L N V P Y I A G I L I I Y F F R G V F Y	
TGAATTAGGT TCTTATGGCT CAAATT (SEQ ID NO:7)	5126
E L G S Y G S N	

FIG. 2a

NFDIETTTFE AMKKHASLLE KISVERSFIE FDKLLLAPYW RKGMLALIDS	50
HAFNYLPCLK NRELQLSAFL SQLDKDFLFE TSEQAWASLI LSMEVEHTKT	100
FLKKWKTSTH FQKDVEHIVD VYRIREQMGL AKEHLYRYGK TIIKQAEGIR	150
KARGLMVDFE KIEQLDSELA IHDRHEIVVN GGTLLIKLGI KPGPQMGDII	200
SQIELAIVLG QLINEEEAIL HFVKQYLM (SEQ ID NO:8)	229

FIG. 2b

MSDFLVDGLT KSVGDKTVFS NVSFIIHSLD RIGIIGVNGT GKTTLLDVIS	50
GELGFDGDRS PFSSANDYKI AYLQEPDFD DSQTILDVTL SSDLREMALI	100
KEYELLNHY EESKQSRLEK VMAEMDSLDA WSIESEVKTV LSKLGITDLQ	150
LSVGELSGGL RRRVQLAQVL LNDADLLLLD EPTNHLDIDT IAWLTNFLKN	200
SKKTVLFITH DRYFLDNVAT RIFELDKAQI TEYQGNYQDY VRLRAEQDER	250
DAASLHKKKQ LYKQELAWMR TQPQARATKQ QARINRFQNL KNDLHQTSDT	300
SDLEMTFETS RIGKKVINFE NVFSYDPKS ILKDFNLLIQ NKDRIGIVGD	350
NGVGKSTLLN LIVQDLQPDS GNVSIGETIR VGYFSQQLHN MDGSKRVINY	400
LQEVADEVKT SVGTTSVTEL LEQFLFPRST HGTQIAKLSG GEKKRLYLLK	450
ILIEKPNVLL LDEPTNDLDI ATLTVLENFL QGFGGPVITV SHDRYFLDKV	500
ANKIIIAFEDN DIREFFGNYT DYLDEKAFNE QNNEVISKKE STKTSREKQS	550
RKRMSYFEKQ EWATIEDDIM ILENTITRIE NDMQTCGSDF TRLSDLQKEL	600
DAKNEALLEK YDRYEYLSEL DT (SEQ ID NO:9)	622

FIG. 2c

MIIRPIIKND DQAVAQLIRO SLRAYDLDKP DTAYSDPHLD HLTSYYEKIE	50
KSGFFVIEER DEIIGCGGFG PLKNLIAEMQ KVYIAERFRG KGLATDLVKM	100
IEVEARKIGY RQLYLETAST LSRATAVYKH MGYCALSQPI ANDQGHTAMD	150
IWMIKDL (SEQ ID NO:10)	157

FIG. 2d

MAYIWSYLR YPNWLWLDLL GAMLFVTVIL GMPTALAGMI DNGVTKGDRT 50
GVYLWTFIMF IFVVLGIIGR ITMAYASSRL TTTMIRDMLRN DMYAKLQEYS 100
HHEYEQIGVS SLVTRMTSDT FVLMQFAEMS LRLGLVTPMV MIFSVVMILI 150
TSPSLAWLVA VAMPLLVGVV LYVAIKTKPL SERQQTMLDK INQYVRENLT 200
GLRVVRAFAR ENFQSQKFQV ANQRYTDST GLFKLTGLTE PLFVQIIIAM 250
IVAIWWFALD PLQRGAIKIG DLVAFIEYSF HALFSFLLFA NLFTMYPRMV 300
VSSHRIREVM DMPISINPNA EGVTDTKLKG HLEFDNVTFA YPGETESPVL 350
HDISFKAKPG ETIAFIGSTG SGKSSLVNLI PRFYDVTLGK ILVDGVDVRD 400
YNLKSLRQKI GFIPQKALLF TGTIGENLKY GKADATIDDL RQAVDISQAK 450
EFIESHQEAF ETHLAEGGSN LSGGQKQRLS IARAVVKDPD LYIFDDSFSA 500
LDYKTDAATLR ARLKEVTGDS TVLIVAQRVG TIMDADQIIV LDEGEIVGRG 550
THAQLIENNA IYREIAESQL KNQNLSEGE (SEQ ID NO:11) 579

FIG. 2e

MRKKSVFLRL WSYLTRYKAT LFLAIFLKVL SSFMSVLEPF ILGLAITELT 50
ANLVDMAKGV SGAELNVPYI AGILIIYFFR GVFYELGSYG SN 92
(SEQ ID NO:12)

FIG. 2f

AATTTGGAAG TGCTCTATCA ACAGTTGAAG TAAAGGAGAT TATTAGTCAA GAAAACATAT 60
 F G S A L S T V E V K E I I S E E N I W
 ---->
 GGTTATATCG GCTCAGTTGC TGCCATTTA CTAGCTACTC ATATTGGAAG TTACCAACTT 120
 L Y R L S C C H F T S Y S Y W K L P T W
 GGTAAGCATC ATATGGGTCT AGCAACAAAG GACAATCAGA TTGCCTATAT TGATGACAGC 180
 M G L A T K D N Q I A Y I D D S
 |---->
 AAAGGTAAGG CAAAAGCCCC TAAAACAAAC AAAACGATGG ATCAAATCAG TGCTGAAGAA 240
 K G K A K A P K T N K T M D Q I S A E E
 GGCATCTCTG CTGAACAGAT CGTAGTCAAA ATTACTGACC AAGGCTATGT GACCTCACAC 300
 G I S A E Q I V V K I T D Q G Y V T S H
 GGTGACCATT ATCATTTTA CAATGGAAA GTTCCTTATG ATGCGATTAT TAGTGAAGAG 360
 G D H Y H F Y N G K V P Y D A I I S E E
 TTGTTGATGA CGGATCCTAA TTACCGTTT AAACAATCAG ACGTTATCAA TGAAATCTTA 420
 L L M T D P N Y R F K Q S D V I N E I L
 |---->
 GACGGTTACG TTATTAAGT CAATGGCAAC TATTATGTT ACCTCAAGCC AGGTAGTAAG 480
 D G Y V I K V N G N Y Y V Y L K P G S K
 CGCAAAACA TTCGAACCAA ACAACAAATT GCTGAGCAAG TAGCCAAAGG AACTAAAGAA 540
 R K N I R T K Q Q I A E Q V A K G T K E
 GCTAAAGAAA AAGGTTAGC TCAAGTGGCC CATCTCAGTA AAGAAGAAGT TGGGGCAGTC 600
 A K E K G L A Q V A H L S K E E V A A V
 AATGAAGCAA AAAGACAAGG ACGCTATACT ACAGACGATG GCTATATTT TAGTCCGACA 660
 N E A K R Q G R Y T T D D G Y I F S P T
 GATATCATTG ATGATTAGG AGATGCTTAT TTAGTACCTC ATGGTAATCA CTATCATTAT 720
 D I I D D L G D A Y L V P H G N H Y H Y
 ATTCCAAAAA AGGATTGTC TCCAAGTGAG CTAGCTGCTG CACAAGCCTA CTGGAGTCAA 780
 I P K K D L S P S E L A A A Q A Y W S Q
 AAACAAGGTC GAGGTGCTAG ACCGTCTGAT TACCGCCCGA CACCAGCCCC AGGTCTAGG 840
 K Q G R G A R P S D Y R P T P A P G R R
 AAAGCCCCAA TTCCTGATGT GACGCCTAAC CCTGGACAAG GTCATCAGCC AGATAACGGT 900
 K A P I P D V T P N P G Q G H Q P D N G
 GGCTATCATC CAGCGCCTCC TAGGCCAAAT GATGCGTCAC AAAACAAACA CCAAAGAGAT 960
 G Y H P A P P R P N D A S Q N K H Q R D
 GAGTTAAAG GAAAAACCTT TAAGGAACCTT TTAGATCAAC TACACCGTCT TGATTTGAAA 1020
 E F K G K T F K E L L D Q L H R L D L K
 TACCGTCATG TGGAAGAAGA TGGGTTGATT TTTGAACCGA CTCAAGTGAT CAAATCAAAC 1080
 Y R H V E E D G L I F E P T Q V I K S N
 GCTTTGGGT ATGTGGTGCC TCATGGAGAT CATTATCATA TTATCCAAAG AAGTCAGTTA 1140
 A F G Y V V P H G D H Y H I I P R S Q L
 TCACCTCTTG AAATGGAATT AGCAGATCGA TACTTAGCTG GCCAAACTGA GGACAATGAC 1200
 S P L E M E L A D R Y L A G Q T E D N D
 TCAGGTTCAAG AGCACTCAA ACCATCAGAT AAAGAAGTGA CACATACCTT TCTTGGTCAT 1260

S	G	S	E	H	S	K	P	S	D	K	E	V	T	H	T	F	L	G	H	
CGCATCAAAG	CTTACGGAAA	AGGCTTAGAT	GGTAAACCAT	ATGATACGAG	TGATGCTTAT	1320														
R	I	K	A	Y	G	K	G	L	D	G	K	P	Y	D	T	S	D	A	Y	
GT	TTT	TAGTA	AAGAATCCAT	TCATTCACTG	GATAAATCAG	GAGTTACAGC	TAAACACGGA	1380												
V	F	S	K	E	S	I	H	S	V	D	K	S	G	V	T	A	K	H	G	
GATCATTTCC	ACTATATAGG	ATTGGAGAA	CTTGAACAAT	ATGAGTTGGA	TGAGGTCGCT	1440														
D	H	F	H	Y	I	G	F	G	E	L	E	Q	Y	E	L	D	E	V	A	
AACTGGGTGA	AAGCAAAAGG	TCAAGCTGAT	GAGCTTGTG	CTGCTTGGGA	TCAGGAACAA	1500														
N	W	V	K	A	K	G	Q	A	D	E	L	A	A	A	L	D	Q	E	Q	
GGCAAAGAAA	AACCACCTT	TGACACTAAA	AAAGTGAGTC	GCAAAGTAAC	AAAAGATGGT	1560														
G	K	E	K	P	L	F	D	T	K	K	V	S	R	K	V	T	K	D	G	
AAAGTGGGCT	ATATGATGCC	AAAAGATGGT	AAGGACTATT	TCTATGCTCG	TGATCAACTT	1620														
K	V	G	Y	M	M	P	K	D	G	K	D	Y	F	Y	A	R	D	Q	L	
GATTTGACTC	AGATTGCCTT	TGCCGAACAA	GAACTAATGC	TTAAAGATAA	GAAGCATTAC	1680														
D	L	T	Q	I	A	F	A	E	Q	E	L	M	L	K	D	K	K	H	Y	
CGTTATGACA	TTGTTGACAC	AGGTATTGAG	CCACGACTTG	CTGTAGATGT	GTCAAGTCTG	1740														
R	Y	D	I	V	D	T	G	I	E	P	R	L	A	V	D	V	S	S	L	
CCGATGCATG	CTGGTAATGC	TACTTACGAT	ACTGGAAGTT	CGTTGTTAT	CCCACATATT	1800														
P	M	H	A	G	N	A	T	Y	D	T	G	S	S	F	V	I	P	H	I	
GATCATATCC	ATGTCGTTCC	GTATTCATGG	TTGACGCGCG	ATCAGATTGC	AACAGTCAAG	1860														
D	H	I	H	V	V	P	Y	S	W	L	T	R	D	Q	I	A	T	V	K	
TATGTGATGC	AACACCCCCGA	AGTTCGTCCG	GATGTATGGT	CTAACGCCAGG	GCATGAAGAG	1920														
Y	V	M	Q	H	P	E	V	R	P	D	V	W	S	K	P	G	H	E	E	
TCAGGTTCCG	TCATCCAAA	TGTTACGCCT	CTTGATAAAC	GTGCTGGTAT	GCCAAACTGG	1980														
S	G	S	V	I	P	N	V	T	P	L	D	K	R	A	G	M	P	N	W	
CAAATTATCC	ATTCTGCTGA	AGAAGTTCAA	AAAGCCCTAG	CAGAAGGTG	TTTTGCAACA	2040														
Q	I	I	H	S	A	E	E	V	Q	K	A	L	A	E	G	R	F	A	T	
CCAGACGGCT	ATATTTTCGA	TCCACGAGAT	GT	TTTGGCCA	AAGAAACTTT	2100														
P	D	G	Y	I	F	D	P	R	D	V	L	A	K	E	T	F	V	W	K	
GATGGCTCCT	TTAGCATCCC	AAGAGCAGAT	GGCAGTTCAT	TGAGAACCAT	TAATAAATCT	2160														
D	G	S	F	S	I	P	R	A	D	G	S	S	L	R	T	I	N	K	S	
GATCTATCCC	AAGCTGAGTG	GCAACAAAGCT	CAAGAGTTAT	TGGCAAAGAA	AAATACTGGT	2220														
D	L	S	Q	A	E	W	Q	Q	A	Q	E	L	L	A	K	K	N	T	G	
GATGCTACTG	ATACGGATAA	ACCCAAAGAA	AAGCAACAGG	CAGATAAGAG	CAATGAAAAC	2280														
D	A	T	D	T	D	K	P	K	E	K	Q	Q	A	D	K	S	N	E	N	
CAACAGCCAA	GTGAAGCCAG	TAAAGAAGAA	AAAGAACATCG	ATGACTTTAT	AGACAGTTA	2340														
Q	Q	P	S	E	A	S	K	E	E	K	E	S	D	D	F	I	D	S	L	
CCAGACTATG	GTCTAGATAG	AGCAACCCCTA	GAAGATCATA	TCAATCAATT	AGCACAAAAAA	2400														
P	D	Y	G	L	D	R	A	T	L	E	D	H	I	N	Q	L	A	Q	K	
GCTAATATCG	ATCCTAAGTA	TCTCATTTC	CAACCAGAAG	GTGTCCAATT	TTATAATAAA	2460														
A	N	I	D	P	K	Y	L	I	F	Q	P	E	G	V	Q	F	Y	N	K	

AATGGTGAAT TGTTAACTTA TGATATCAAG ACACCAAC AAATAAACCC TTAACCAAAA 2520
 N G E L V T Y D I K T L Q Q I N P

 GAAGATCTCA TTGTTAAAGC ACTGCTTGT CAAAGCAAGT TACGGTGATT TTGAAGTCAT 2580

 TCTATGTAAC GAGTAGTGAT AAAAGTTGGA TAATAGCGGT TTTCTTTGC AAAGAAATGG 2640

 TATCCATGTT AGAATAGTAA AAAAAGAGGA GGATTCTTGG ACTAATGTCA AATAAGTAGA 2700

 CAGAAAAC TGTTATTTA TTGCGTTAAA ATAATTTCT TCTTCTGAT TAGGGGTTAG 2760
 .K I A N F Y N E E K Q N P T L

 TCCTAGATTA GCGTATGTG GGGTGTAAATT GTTATAAAAAA TTCTCAATGT ATTCAAAGCA 2820
 G L N A T H P N Y N N Y F N E I Y E F C

 GTCTAATTGA ACCTGTTGA TATTTGATA ATGTTTCGG TTGATTGTC TATGCTTTAA 2880
 D L Q V Q K I N Q Y H K R N I Q R H K L

 ATACTGAAA AATGCTTCAG TTACGGCATT ATCATAAGGA TATCCAGGAT TAGAAAAAGA 2940
 Y K F F A E T V A N D Y P Y G P N S F S

 ATGCATGATA TTGGCACTGC ACCCTAATAG TGAGACGCAA GAAAAACACT TTTAGGCAAT 3000
 H M A I
 <----|
 CAGTTTCTG TACTGTACAG GCGACTGGTC GTTTAATCTC TGTTGAATT TAGTTCTATT 3060
 L K R Y Q V P S Q D N L R Q Q I R T E N

 ATAAAATGTA ATGTAATTT TAACAATATT TGTTATACTA TCTTGTGT ATTTCTCCT 3120
 Y F T I Y N K V I N T I S D K N Y K R R

 ATTATGGAAA TAAAAGGTTT CAGTCTTAG GACGGTGTGA AACCAATTCAA TACAGGCATT 3180
 N H F Y F T E T K L V T H F W E I C A N

 ATCTGCAGGT GTTCCTTTTC GAGACATTGA GCGGATAATG TCTTTTCCG TGCAAGCCTG 3240
 D A P T G K R S M S R I I D K E T C A Q

 GTAGTAAGCC ATAGAAGTAT ACACTGAGCC TTGGTCACTG TGTAAGATTG CTCTTTATT 3300
 Y Y A M
 <----|
 TAGGCAATT TAACTGATTA AGGGTGTCTA GTACAAAATC CGTGTCTGA CAATCTGAGA 3360
 K P L K L Q N L T D L V F D T D Q C D S

 TAGTGTAAAGC TATAATTCT CGGTTATAGA GATTCTAAAT TGATGAGAGA TACAATTAC 3420
 I T Y A I I E R N Y L N M I S S L Y L K

 AGTTACCGAA ATATAAGGTAG GAAATATCTG TTACGAGCTT TTCTTAGGC TTATCGGCAT 3480
 C N G F Y L Y T I D T V L K E K P K D A

 GGAAATCCCC ACTCAATTAA TTATCTGTTA AATAATAAGC TTTACCCAAA TTGGGAACCTT 3540
 H G D R S L K N D T L Y Y A K G L N P V

 TCTTGGTACG TGTCCGACAA AGCCAGCCAT TATTTTCAT GATACGATAG ACTTTCTTG 3600
 K K T R T R C L W G N N K M I R Y V K K

 TATTAACAGT CAATCCGTGG ATTTTTTGAA GCAATCGTGT AATGGTACGA TAGCCATAAA 3660
 T N V T L G H I K K L L R T I T R Y G Y

 TAAAGTGATT CTCCATACAG AGCTGTTCAA TTAATTCAAT AAGGTACATCT TTTTTGCGG 3720
 I F H N E M
 <----|

CTTCTCATAAC TCCTTTTCC AACGGTAATA GGTCGACCGC TTGACCTAA AACAGTCTAG 3780
 AATGAAAACCT ATCGGGTAGT TGTTTTATA GTCTTCCACA AGCTTGATAA GACTTACTTT 3840
 ATCGATTTCC TTATCAAAGCC TCGATACTTT TTTAAGAGGT CAACCTGTAA TTGTAATTGT 3900
 I S K R I L G R Y K K L L D V Q L Q L Q
 TCCACTTCAG ACAGATGTTCA AAGCCTTAA CCGTAGGTAT ATTGCTTGCC AACACCTTGA 3960
 E V E S L H E L G K G Y T Y Q K G V G Q
 TGAAAACGAT AAAGCTCCTC GTTTCTGTAC CATTTCATCC AAGTATAGAT TTGACTATTAA 4020
 H F R Y L E E N E Y W K M W T Y I Q S N
 TTTTGATGC CTAAAGTCTC CATAATAACT CTGTTAGACT TGCCCTGCTTT CTTCATATCG 4080
 N K I G L T E M I V R N S K G A K K M D
 ATGCAAGCCA GCTTAGTTTC CCATGAATAT GCTTTTTAA CCATAATAAA ACATTCCGT 4140
 I C A L K T E W S Y A K K V M
 <----|
 TTCTAGTTA CTAAATTCA ACAGGAGTGT TTTCTTTG TCTCATTAA GGGATTCACT 4200
 GCCTATTGTT GTCATCAATT ATTTTCTAA ATTCCCCGGA CTTAAATTGT GACCCTTGGT 4260
 CGGAATGAAA GAGAAGTGT CTTCAATCT TTCTTTATT AAGTAAAAG GCAACACTTT 4320
 TCTGTACAAC ATTTATAAAAG TGTTTTCTA GGCAATTAAT CTTTAGTCA TTGGTGTGGT 4380
 . A I L R K T M P T Q
 GTAGTTGAGA CTACCATGAA TGCGGTGGTA ATTCCACCAA TGAACATAGT CTTAGTCTT 4440
 Y N L S G H I R H Y N W W H V Y D K T K
 AAGAGCTAGT TCTTCCAGCA ATTGAAAGGT TTCTTGATAA ACAAAATTCAA TTTGAAAGC 4500
 L A L E E L L Q F T E Q Y V F E I K F A
 ACGATACGTA CTTTCAGCTA CGGCATTGTC ATAAGGATAA CCAGCCTGAC TAAGCGAACG 4560
 R Y T S E A V A N D Y P Y G A Q S L S R
 TGTGATTCCA AAGGCTTCCA ATATTCATC AATTAACGTAA TTATCAAACCT CTTGCCACG 4620
 T I G F A E L I E D I L Q N D F E K G R
 ATCTGAATGG AACATCTTGA CTTGGTCAG GGCATAAGGG ATGCTTGTA TGGCTTGCTT 4680
 D S H F M K V K T L A Y P I S Q I A Q K
 AACGAGTTCA GCGGTCTTGT GCCAACCAAG AGACAGGCCG ATGATTTCAC GGTGTATAG 4740
 V L E A T K H W G L S L G I I E R N Y L
 GTCATGATG AGGCAACAT AAGCCCAACG ATTGCCTACA CGAACATAGG TTAAGTCAGT 4800
 D I I L C V Y A W R N G V R V Y T L D T
 GACTAAGGCT TGTAGTGGTC TTTCTTGCTT AAATTGCCTG TCTAAGTGGT TGGGAATAGG 4860
 V L A Q L P R E Q K F Q R D L H N P I P
 GGCTTCATTC TTGCCTCTAG AATGTGGTT GAAGGTGGCT TTCTGATAAA CAGAAACCAA 4920
 A E N K G R S H P K F T A K Q Y V S V L
 ATTGAGTCGC TTCATAATGC GTGAAATCCG ACGACGTGAA AGTGTGATAC CTTCGTTATT 4980
 N L R K M I R R I R R R S L T I G E N N
 CAAGCATATT TTGATTTTC TGGATCCGTA TCTAGACTCG CTATCGAGAA AAATTCTTTT 5040
 L C I K I K R S G Y R S E S D L F I R K

AATAGTTCT	TCAAACTCCG	TTTCAGATAC	TGACTCCACG	GCTTGATAGT	AATAACTTGA	5100
I	T	E	E	F	E	
			T	E	S	V
				S	E	V
				A	Q	Y
					Y	Y
					S	S
GTGTGGCATA TTCAGCCAGC GACACATCTT TGAAATGCTG TATTATCCT TATTAGCAGT						5160
H	P	M	N	L	W	R
				C	M	K
				S	I	S
					Y	K
					D	D
					K	N
					A	T
GATTATTCCT CTTTTGTGC CATAATCACC GCTGCTTGCT TTAGGATATC TAATT						5215
I	I	E	R	K	T	
				G	Y	D
				D	G	S
				S	S	A
				A	K	P
					Y	Y
					R	I

(SEQ ID NO:13)

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FIG. 3a

FGSALSTVEV KEIISEENIW LYRLSCCHFT SYSYWKLPTW

40

(SEQ ID NO:14)

FIG. 3b

MGLATKDNQI	AYIDDSKGKA	KAPKTNKTMD	QISAEEGISA	EQIVVKITDQ	50
GYVTSHGDHY	HFYNGKVPYD	AIISEELLMT	DPNYRFKQSD	VINEILDGYV	100
IKVNGNYYVY	LKPGSKRKNI	RTKQQIAEQV	AKGTKEAKEK	GLAQVAHLSK	150
EEVAAVNEAK	RQGRYTTDDG	YIFSPTDIID	DLGDAYLVPH	GNHYHYIPKK	200
DLSPSELAAA	QAYWSQKQGR	GARPSDYRPT	PAPGRRKAPI	PDVTPNPGQG	250
HQPDNGGYHP	APPRPNDASQ	NKHQRDEFKG	KTFKELLDQL	HRLDLKYRHV	300
EEDGLIFEPT	QVIKSNAFGY	VVPHGDHYHI	IPRSQLSPLE	MELADRYLAG	350
QTEDNDSGSE	HSKPSDKEVT	HTFLGHRIKA	YGKGLDGKPY	DTSDAYVFSK	400
ESIHSVVDKSG	VTAKHGDHFH	YIGFGELEQY	ELDEVANWVK	AKGQADELAA	450
ALDQEQQKEK	PLFDTKKVSR	KVTKDGKVGY	MMPKDGDYF	YARDQLDLTQ	500
IAFAEQELML	KDKKHYRYDI	VDTGIEPRLA	VDVSSLPMHA	GNATYDTGSS	550
FVIPHIDHIH	VVPYSWLTRD	QIATVKYVMQ	HPEVRPDWWS	KPGHEESGSV	600
IPNVTPLDKR	AGMPNWQIIH	SAEEVQKALA	EGRFATPDGY	IFDPRDVLA	650
ETFVWKDGGSF	SIPRADGSSL	RTINKSDLSQ	AEWQQAQELL	AKKNTGDA	700
TDKPKEKQQA	DKSNENQQPS	EASKEEKESD	DFIDSLPDYG	LDRATLEDHI	750
NQLAQKANID	PKYLIFQPEG	VQFYNKNGEL	VTYDIKTLQQ	INP	793

(SEQ ID NO:15)

FIG. 3c

MTDPNYRFKQ SDVINEILDG YVIKVNGNYY VYLPKGSKRK NIRTQQIAE	50
QVAKGTKEAK EKGLAQVAHL SKEEVAAVNE AKRQGRYTTD DGYIFSPTDI	100
IDDLGDAYLV PHGNHYHYIP KKDLSPSELA AAQAYWSQKQ GRGARPSDYL	150
PTPAPGRKA PIPDVTPNPG QGHQPDNGGY HPAPPRNDA SQNKHQRDEF	200
KGKTFKELLD QLHRLDLKYR HVEEDGLIFE PTQVIKSNAF GYVVPHGDHY	250
HIIPRSQLSP LEMELADRYL AGQTEDNDSG SEHSKPSDKE VTHTFLGHRI	300
KAYGKGLDGK PYDTS DAYVF SKESIHSVDK SGVTAKHGDH FHYIGFGELE	350
QYELDEVANW VKAKGQADEL AAALDQEQQK EKPLFDKKV SRKVTKDGKV	400
GYMMPKDGKD YFYARDQLD TQIAFAEQEL MLKDKKHYRY DIVDTGIEPR	450
LAADVSSLPM HAGNATYDTG SSFVIPHIDH IHVVPYSWLT RDQIATVKYV	500
MQHPEVRPDV WSKPGHEESG SVIPNVTPLD KRAMGPNWQI IHSAAEVQKA	550
LAEGRFATPD GYIFDPRDVL AKETFVWKDG SFSIPRADGS SLRTINKSDL	600
SQAEWQQAQE LLAKKNTGDA TDTDKPKEKQ QADKSNNENQQ PSEASKEEKE	650
SDDFIDSLPD YGLDRATLED HINQLAQKAN IDPKYLIFQP EGVQFYNKNG	700
ELVTYDIKTL QQINP (SEQ ID NO:16)	715

FIG. 3d

MHSFSNPGYP YDNAVTEAFF KYLKHRQINR KHYQNIKQVQ LDCFEYIENF	50
YNNYNPHTAN LGLTPNQKEE NYFNAIK (SEQ ID NO:17)	77

FIG. 3e

MAYYQACTEK DIIRSMSRKG TPADNACIEW FHTVLKTETF YFHNRKYNK	50
DSITNIVKNY ITFYNETRIQ QRLNDQSPVQ YRKLIA (SEQ ID NO:18)	86

FIG. 3f

MENHFIYGYR TITRLLKKIH GLTVNTKKVY RIMKNNGWLC RTRTKKVPNL	50
GKAYYLTDNK LSRDFHADKP KEKLVTDITY LYFGNCKLYL SSIMNLYNRE	100
IIAYTISDCQ DTDFVLDTLN QLKLPK (SEQ ID NO:19)	126

FIG. 3g

MVKKAYSWET KLACIDMKKA GKSNRVIMET LGIKNNSQIY TWMKWYENEE 50
 LYRFHQGVGK QTYGKGLEH LSEVEQLQLQ VDLLKKYRGL IRKSIK 96
 (SEQ ID NO:20)

FIG. 3h

IRYPKASSGD YGTKREIITA NKDKYSISKM CRWLNMIPHSS YYYQAVESVS 50
 ETEFEETIKR IFILDSESRYG SRKIKICLNN EGITLSRRRI RRIMKRLNLV 100
 SVYQKATFKP HSRGKNEAPI PNHLDRQFKQ ERPLQALVTD LTYVRVGNRW 150
 AYVCLIIDL NREIIGLSLG WHKTAELVKQ AIQSIPYALT KVKMFHSDRG 200
 KEFDNQLIDE ILEAFGITRS LSQAGYPYDN AVAESTYRAF KIEFVYQETF 250
 QLLEELALKT KDYVHWWNYH RIHGSILNYQT PMTKRLIA (SEQ ID NO:21) 288

FIG. 3i

AATTGAAAG CAGAATTATC TGTAGAAGAT GAGCAATATA CAGCAACAGT TTATGGTAA 60
 N L K A E L S V E D E Q Y T A T V Y G K
 ---->
 TCTGCTCATG GTTCAACACC ACAAGAAGGT GTTAATGGGG CGACTTATTT AGCTCTTTAT 120
 S A H G S T P Q E G V N G A T Y L A L Y
 CTAAGTCAAT TTGATTTGA AGGTCTGCT CGTGCTTCT TAGATGTTAC AGCCAACATT 180
 L S Q F D F E G P A R A F L D V T A N I
 ATTCACGAAG ACTTCTCAGG TGAAAAACTT GGAGTAGCTT ATGAAGATGA CTGTATGGGA 240
 I H E D F S G E K L G V A Y E D D C M G
 CCATTGAGCA TGAATGCAGG TGTCTTCCAG TTTGATGAAA CTAATGATGA TAATACTATC 300
 P L S M N A G V F Q F D E T N D D N T I
 GCTCTTAATT TCCGTACCC ACAAGGGACA GATGCTAAA CTATCCAAAC TAAGCTTGAG 360
 A L N F R Y P Q G T D A K T I Q T K L E
 AAACCTAACG GAGTTGAAAA AGTGACTCTT TCTGACCATG AACACACACC ACACATATGTA 420
 K L N G V E K V T L S D H E H T P H Y V
 CCTATGGACG ATGAATTAGT ATCAACCTTA CTAGCTGTCT ATGAAAAGCA AACTGGTCTT 480
 P M D D E L V S T L L A V Y E K Q T G L
 AAAGGACATG AACAGGTTAT TGGTGGTGGG ACATTTGGTC GCTTACTTGA ACGGGGTGTT 540
 K G H E Q V I G G G T F G R L L E R G V
 GCATACGGTG CCATGTTCCC AGGAGATGAA AACACTATGC ATCAAGCTAA TGAGTACATG 600
 A Y G A M F P G D E N T M H Q A N E Y M
 CCTTTAGAAA ATATTTCCG TTCGGCTGCT ATCTACGCAG AAGCTATCTA TGAATTAATC 660

P L E N I F R S A A I Y A E A I Y E L I
 AAATAAAATA ATCCTTAAAC TAAATATGTG ATCAATGATA AAGGGTGGTG AAGACATGAA 720
 K .

AGTGTCTTG CCTCTTTCA TAAGGTTAGA TTTGGAGACT TTATGACTGA CTTGGAAAAA 780
 M T D L E K
 |---->
 ATTATTAAGA CAATAAAAAG TGATTCACAG AATCAAATT ATACAGAAA TGGTATTGAT 840
 I I K A I K S D S Q N Q N Y T E N G I D

CCTTGTTTG CTGCTCCTAA AACAGCTAGG ATCAATATTG TTGGCCAAGC ACCTGGTTA 900
 P L F A A P K T A R I N I V G Q A P G L

AAAACTCAAG AAGCAAGACT CTATTGGAAA GATAAATCTG GAGATCGTCT ACGCCAGTGG 960
 K T Q E A R L Y W K D K S G D R L R Q W

CTTGGAGTTG ATGAAGAGAC ATTTTACCAT TCTGGAAAAT TTGCTGTTT ACCTTTAGAT 1020
 L G V D E E T F Y H S G K F A V L P L D

TTTTATTACC CAGGCAAAGG AAAATCAGGA GATTTACCC CTAGAAAAGG TTTGCGGAG 1080
 F Y Y P G K G K S G D L P P R K G F A E

AAATGGCACC CTCTTATTT AAAAGAAAATG CCTAATGTC AATTGACCTT GCTAGTTGGT 1140
 K W H P L I L K E M P N V Q L T L L V G

CAGTATGCTC AGAAATATTA TCTTGGAAAGC TCCGCACATA AAAATCTAAC AGAAACAGTT 1200
 Q Y A Q K Y Y L G S S A H K N L T E T V

AAAGCTTACA AAGACTATCT ACCCGATTAT TTACCCCTGG TTCACCCATC ACCCGCAAAT 1260
 K A Y K D Y L P D Y L P L V H P S P R N

CAAATTTGGC TAAAGAAGAA TCCATGGTTT GAAAAGATC TAATCGTGA TTTACAAAAG 1320
 Q I W L K K N P W F E K D L I V D L Q K

ATAGTAGCAG ATATTTAAA AGATTAAGGA TAGGAGTTGG TATGAGAGAT AATCATCTAC 1380
 I V A D I L K D . M R D N H L H
 |---->

ACACGTATTT TTCCTATGAT TGTCAAACGG CATTGAGGA CTATATTAAT GGTTTACAG 1440
 T Y F S Y D C Q T A F E D Y I N G F T G

GTGAATTTAT CACGACAGAA CATTGATT TATCAAATCC TTACACCGGT CAAGACGATG 1500
 E F I T T E H F D L S N P Y T G Q D D V

TTCCTGATTA TAGTGCCTAT TGTCAAAAAA TAGATTATCT TAATCAGAAA TATGGAAATC 1560
 P D Y S A Y C Q K I D Y L N Q K Y G N R

GATTTAAAAA AGGAATTGAA ATCGGTTATT TTAAAGATAG GGAATCAGAT ATTTAGATT 1620
 F K K G I E I G Y F K D R E S D I L D Y

ATTTAAAAAA TAAAGAATTGATT TATTGTCAT CCATCATAAT GGTAGGTATG 1680
 L K N K E F D L K L L S I H H N G R Y D

ATTATCTGCA AGAAGAAGCT CTGAAAGTAC CAACAAAGGG AGCTTTAGC AGATTACTTT 1740
 Y L Q E E A L K V P T K G A F S R L L .

AATCGTATGG AATTTGCCAT AGGCCGTGTG GAAGCGCACG TTTAGCTCA CTTTGATTAT 1800

GGTTTCGTA AGTTAAACTT AGATGTAGAA GATTTAAAAC CGTTGAAAC GCAATTGAAG 1860

CGCATTTCGA TAAAGATGTT ATCTAAGGGG TTAGCTTTG AACTAAATAC CAAATCCCTT 1920

TATCTATATG GGAATGAAAA ACTTTATCGC TATGCTTAG AGATACTCAA ACAGCTTGGT 1980
 TGTAACAAAT ACTCTATAGG CTCTGACGGT CATATTCTG AACATTTTG TTATGAATT 2040
 GATAGACTTC AAGGTCTGCT AAAGGACTAT CAAATTGATG AAAATCATT GATATGAGGA 2100
 AATTTTGAT AAAAAAGCTA GGCAATATTG CTTAGCTTT TTGTAATGCT ATTGATAGTT 2160
 TTAGTGAAAA TTTCAAAAAA ATAAAGAAAT CATTACTTG TTGCAAGCGC TTGCGTAAAT 2220
 TGTTATGATT TTATTGGTAA CAATTCACTTA AAAAAGGAGA ATGATATGAA AAGAAAAGAC 2280
 M K R K D
 |---->
 TTATTTGGTG ATAAACAAAC TCAATACACG ATTAGAAAGT TAAGTGTGG AGTAGCTTCA 2340
 L F G D K Q T Q Y T I R K L S V G V A S
 GTTACAACAG GGGTATGTAT TTTCTTCAT AGTCCACAGG TATTTGCTGA AGAAGTAAGT 2400
 V T T G V C I F L H S P Q V F A E E V S
 GTTTCTCCTG CAACTACAGC GATTGCAGAG TCGAATATTAA ATCAGGTTGA CAACCAACAA 2460
 V S P A T T A I A E S N I N Q V D N Q Q
 TCTACTAATT TAAAAGATGA CATAAACTCA AACTCTGAGA CGGTTGTGAC ACCCTCAGAT 2520
 S T N L K D D I N S N S E T V V T P S D
 ATGCCGGATA CCAAGCAATT AGTATCAGAT GAAACTGACA CTCAAAAGGG AGTGACAGAG 2580
 M P D T K Q L V S D E T D T Q K G V T E
 CCGGATAAGG CGACAAGCCT GCTTGAAGAA AATAAAGGTC CTGTTTCAGA TAAAATACC 2640
 P D K A T S L L E E N K G P V S D K N T
 TTAGATTTAA AAGTAGCACC ATCTACATTG CAAAATACTC CCGACAAAAC TTCTCAAGCT 2700
 L D L K V A P S T L Q N T P D K T S Q A
 ATAGGTGCTC CAAGCCCTAC CTTGAAAGTA GCTAATCAAG CTCCACGGAT TGAAAATGGT 2760
 I G A P S P T L K V A N Q A P R I E N G
 TACTTTAGGC TACATCTTAA AGAATTGCCT CAAGGTCATC CTGTAGAAAG CACTGGACTT 2820
 Y F R L H L K E L P Q G H P V E S T G L
 TGGATATGGG GAGATGTTGA TCAACCGTCT AGTAATTGGC CAAATGGTGC TATCCCTATG 2880
 W I W G D V D Q P S S N W P N G A I P M
 ACTGATGCTA AGAAAGATGA TTACGGTTAT TATGTTGATT TTAAATTATC TGAAAACAA 2940
 T D A K K D D Y G Y Y V D F K L S E K Q
 CGAAAACAAA TATCTTTTT AATTAATAAC AAAGCAGGGA CAAATTAAAG CGGCGATCAT 3000
 R K Q I S F L I N N K A G T N L S G D H
 CATATTCCAT TATTACGACC TGAGATGAAC CAAGTTGGA TTGATGAAAA GTACGGTATA 3060
 H I P L L R P E M N Q V W I D E K Y G I
 CATACTTATC AACCCCTCAA AGAAGGGTAT GTCCGTATTA ACTATTGAG TTCCCTCTAGT 3120
 H T Y Q P L K E G Y V R I N Y L S S S S
 AACTATGACC ACTTATCAGC ATGGCTCTT AAAGATGTTG CAACCCCYTC AACAACTTGG 3180
 N Y D H L S A W L F K D V A T P S T T W
 CCAGATGGTA GTAATTTGT GAATCAAGGA CTATATGGAA GGTATATTGA TGTATCACTA 3240
 P D G S N F V N Q G L Y G R Y I D V S L

AAAACTAACG CCAAAGAGAT TGGTTTCTA ATCTTAGATG AAAGTAAGAC AGGAGATGCA 3300
 K T N A K E I G F L I L D E S K T G D A

 GTGAAAGTTC AACCCAACGA CTATGTTTT AGAGATTTAG CTAACCATAA CCAAATTTT 3360
 V K V Q P N D Y V F R D L A N H N Q I F

 GTAAAAGATA AGGATCCAAA GGTTTATAAT AATCCTTATT ACATTGATCA AGTGCAGCTA 3420
 V K D K D P K V Y N N P Y Y I D Q V Q L

 AAGGATGCC AACAAATTGA TTTAACAAAGT ATTCAAGCAA GTTTACAAC TCTAGATGGG 3480
 K D A Q Q I D L T S I Q A S F T T L D G

 GTAGATAAAA CTGAAATTTT AAAAGAATTG AAAGTGAATG ATAAAAATCA AAATGCTATA 3540
 V D K T E I L K E L K V T D K N Q N A I

 CAAATTCCTG ATATCACTCT CGATACTAGT AAATCTCTT TAATAATCAA AGGCGACTTT 3600
 Q I S D I T L D T S K S L L I I K G D F

 AATCCTAAAC AAGGTCAATT CAACATATCT TATAATGGTA ACAATGTCAT GACAAGGCAA 3660
 N P K Q G H F N I S Y N G N N V M T R Q

 TCTTGGGAAT TTAAAGACCA ACTTTATGCT TATACTGGAA ATTTAGGTGC AGTTCTCAAT 3720
 S W E F K D Q L Y A Y S G N L G A V L N

 CAAGATGGTT CAAAAGTTGA AGCCAGCCTC TGGTCACCGA GTGCTGATAG TGTCACTATG 3780
 Q D G S K V E A S L W S P S A D S V T M

 ATTATTTATG ACAAAAGATAA CCAAAACAGG GTTGTAGCGA CTACCCCCCT TGTGAAAAAT 3840
 I I Y D K D N Q N R V V A T T P L V K N

 AATAAAGGTG TTTGGCAGAC GATACTTGAT ACTAAATTAG GTATTAAAAA CTATACTGGT 3900
 N K G V W Q T I L D T K L G I K N Y T G

 TACTATTATC TTTACGAAAT AAAAAGAGGT AAGGATAAGG TTAAGATTT AGATCCTTAT 3960
 Y Y Y L Y E I K R G K D K V K I L D P Y

 GCAAAGTCAT TAGCAGAGTG GGATAGTAAT ACTGTTAATG ATGATATTAA AACGGCTAAA 4020
 A K S L A E W D S N T V N D D I K T A K

 GCAGTTTG TAAATCCAAG TCAACTTGA CCTCAAAATT TAAGTTTGC TAAAATTGCT 4080
 A A F V N P S Q L G P Q N L S F A K I A

 AATTTAAAG GAAGACAAGA TGCTGTTATA TACGAAGCAC ATGTAAGAGA CTTCACTTCT 4140
 N F K G R Q D A V I Y E A H V R D F T S

 GATCGATCTT TGGATGGAAA ATTTAAAAAT CAATTTGGTA CCTTTGCAGC CTTTCAGAG 4200
 D R S L D G K L K N Q F G T F A A F S E

 AAACTAGATT ATTTACAGAA ATTAGGAGTT ACACACATTC AGCTTTTACCG GGTATTGAGT 4260
 K L D Y L Q K L G V T H I Q L L P V L S

 TATTTTATG TTAATGAAAT GGATAAGTCA CGCTAACAG CTTACACTTC CTCAGACAAT 4320
 Y F Y V N E M D K S R S T A Y T S S D N

 AATTACAATT GGGGCTATGA CCCACAGAGC TATTTGCTC TTTCTGGGAT GTATTCAAG 4380
 N Y N W G Y D P Q S Y F A L S G M Y S E

 AAACCAAAAG ATCCATCAGC ACGTATGCC GAATTAAAAC AATTAATACA TGATATTCA 4440
 K P K D P S A R I A E L K Q L I H D I H

AACACGTGGCA TGGGGGTTAT ACTTGATGTC GTCTATAATC ACACTGCAAA AACTTATCTC 4500
 K R G M G V I L D V V Y N H T A K T Y L

 TTTGAGGATA TAGAACCTAA TTATTATCAC TTTATGAATG AAGATGGTTC ACCAAGAGAA 4560
 F E D I E P N Y Y H F M N E D G S P R E

 AGTTTTGGAG GGGGACGTTT AGGAACCACT CATGCAATGA GTCGTCGTGT TTTGGTTGAT 4620
 S F G G G R L G T T H A M S R R V L V D

 TCCATTAAT ATCTTACAAG TGAATTTAAA GTTGATGGTT TCCGTTTG A TATGATGGGA 4680
 S I K Y L T S E F K V D G F R F D M M G

 GATCATGATG CGGCTGCGAT TGAATTAGCT TATAAAAGAAG CTAAAGCTAT TAATCCTAAT 4740
 D H D A A A I E L A Y K E A K A I N P N

 ATGATTATGA TTGGTGAGGG CTGGAGAAC A TTCCAAGGCG ATCAAGGTCA GCCGGTTAAA 4800
 M I M I G E G W R T F Q G D Q G Q P V K

 CCAGCTGACC AAGATTGGAT GAAGTCAAC C GATACAGTTG GCGTCTTTTC AGATGATATT 4860
 P A D Q D W M K S T D T V G V F S D D I

 CGTAATAGCT TGAAATCTGG TTTCCAAAT GAAGGTACTC CAGCTTCAT CACAGGTGGC 4920
 R N S L K S G F P N E G T P A F I T G G

 CCACAATCTT TACAAGGTAT TTTAAAAAT ATCAAAGCAC AACCTGGAA TTTGAAGCA 4980
 P Q S L Q G I F K N I K A Q P G N F E A

 GATTCGCCAG GAGATGTGGT GCAGTATAATT GCTGCACATG ATAACCTTAC CTTGCATGAT 5040
 D S P G D V V Q Y I A A H D N L T L H D

 GTGATTGC AATCAATT (SEQ ID NO:22) 5058
 V I A K S I .

FIG. 4a

NLKAELSVED EQYTATVY GK SAHGSTPQEG VNGATYLALY LSQFD FEGPA 50
 RAFLDVTANI IHEDFSGEKL GVAYEDDCMG PLSMNAGVFQ FDETNDNTI 100
 ALNFRYPQGT DAKTIQTKLE KLNGVEKVTL SDHEHTPHYV PMDDELVSTL 150
 LAVYEKQTGL KGHEQVIGGG TFGRLLERGV AYGAMFPGDE NTMHQANEYM 200
 PLENIFRSAA IYAEAIYELI K (SEQ ID NO:23) 221

FIG. 4b

MTDLEKIIKA IKSDSQNQNY TENGIDPLFA APKATARINIV GQAPGLKTQE 50
 ARLYWKDKSG DRLRQWLGV D E E T F Y H S G K F A V L P L D F Y Y P G K G K S G D L P P 100
 RKGFAEKWHP LILKEMPNVQ L T L L V G Q Y A Q K Y Y L G S S A H K N L T E T V K A Y K 150
 D Y L P D Y L P L V H P S P R N Q I W L K K N P W F E K D L I V D L Q K I V A D I L K D 194
 (SEQ ID NO:24)

FIG. 4c

MRDNHLHTYF SYDCQTAFED YINGFTGEFI TTEHFDSLNP YTGQDDVPDY	50
SAYCQKIDYL NQKYGNRFKK GIEIGYFKDR ESDILDYLKN KEFDLKLLSI	100
HHNGRYDYLQ EEALKVPTKG AFSRLL (SEQ ID NO:25)	126

FIG. 4d

MKRKDLFGDK QTQYTIRKLS VGVASVTTGV CIFLHSPQVF AEEVSVSPAT	50
TAIAESNINQ VDNQQSTNLK DDINSNSETV VTPSDMPDTK QLVSDETDTQ	100
KGVTEPDKAT SLLEENKGPV SDKNTLDLK V APSTLQNTPD KTSQAIGAPS	150
PTLKVANQAP RIENGYFRLH LKELPQGHPV ESTGLWIWGD VDQPSSNWPN	200
GAIPMTDAKK DDYGYYVDFK LSEKQRKQIS FLINNKAGTN LSGDHHIPLL	250
RPEMNQVWID EKYGIHTYQP LKEGYVRINY LSSSSNYDHL SAWLFKDVTAT	300
PSTTWPDGSN FVNQGLYGRY IDVSLKTNAK EIGFLILDES KTGDAVKVQP	350
NDYVFRDLAN HNQIFVKDKD PKVYNNPYYI DQVQLKDAQQ IDLTSIQASF	400
TTLDGVDKTE ILKELKVTDK NQNQAIQISDI TLDTSKSLLI IKGDFNPKQG	450
HFNISYNGNN VMTRQSWEFK DQLYAYSGNL GAVLNQDGSK VEASLWSPSA	500
DSVTMIIYDK DNQNRVVATT PLVKNNKGW QTILDTKLGI KNYTGYYYLY	550
EIKRGKDKVK ILDPYAKSLA EWDSNTVNDD IKTAKAAFVN PSQLGPQNLS	600
FAKIANFKGR QDAVIYEAHV RDFTSDRSLD GKLKNQFGTF AAFSEKLDYL	650
QKLGVTHIQL LPVLSYFYVN EMDKSRSTAY TSSDNNYNWG YDPQSYFALS	700
GMYSEKPKDP SARIAELKQL IHDIHKRGMG VI LDVVVNHT AKTYLFEDIE	750
PNYYHFMNED GSPRESFGGG RLGTTHAMSR RVLVDSIKYL TSEFKVDGFR	800
FDMMGDHDAA AIELAYKEAK AINPNMIMIG EGWRTFQGDQ GQPVKPADQD	850
WMKSTDVTGV FSDDIRNSLK SGFPNEGTPA FITGGPQSLQ GIFKNIKAQP	900
GNFEADSPGD VVQYIAAHDN LTLHDVIAKS I (SEQ ID NO:26)	931

FIG. 4e

AATTCAAAGT TTGACAGAAG GTCAACTTCG TTCTGATATC CCTGAGTTCC GTGCTGGTGA 60
 I Q S L T E G Q L R S D I P E F R A G D
 ---->
 TACTGTACGT GTTCACGCTA AAGTTGTTGA AGGTACTCGC GAACGTATTG AGATCTTG 120
 T V R V H A K V V E G T R E R I Q I F E
 AGGTGTTGTT ATCTCACGTA AAGGTCAAGG AATCTCAGAA ATGTACACAG TACGTAAAAT 180
 G V V I S R K G Q G I S E M Y T V R K I
 TTCTGGTGGT ATCGGTGTTAG AGCGTACATT CCCAATTCAC ACTCCTCGTG TTGATAAAAAT 240
 S G G I G V E R T F P I H T P R V D K I
 CGAAGTTGTT CGTTATGGTA AAGTACGTCG TGCTAAACTT TACTACTTAC GCGCATTGCA 300
 E V V R Y G K V R R A K L Y Y L R A L Q
 AGGTAAAGCT GCACGTATTA AAGAAATCCG TCGTTAATTT TGATGATCAG ATTTTAAAAAA 360
 TGCTTGGTTG TTTGAGGATA GTAACTATGT TTTAAAATG GACAACCAAG ACGTAAAAAA 420
 TCTGCCTGTG GGCAGTTTT TTACTAGGTC CCCTTAGTTC AATGGATATA ACAACTCCCT 480
 . H I Y C S G
 CCTAAGGAGT AATTGCTGGT TCGATTCCGG CAGGGGACAT ATTCAATTGCA TGAAATAGC 540
 G L S Y N S T R N R C P V Y E N C T F L
 GGTTTAGAGC TATTTTGCCC CAAATTCTC TGATTAAGTT TATCGTTCCCT ATCTTTTGT 600
 P K S S N Q G L N R Q N L K D N R D K Q
 TCTTGTAATT GATGTGCGTA AACTCTAAA GTGATATTAA ATTCTCGTG ATCTAAAATC 660
 E Q L Q H A Y V E L T I N L N E H D L V
 TGAGAGATGG AAATTAGATA GCTTGCATAT GSTATGCCTGA GAGAGTGCAC TCGTACCTCG 720
 Q S I S I L Y S A F T H R L S H V R V E
 CGACCAAGTTA TTTTCGGAT AGTTTATTG ACTGCATTAT TTGAAAGTTT GTCGAATAAT 780
 R G T I K R I T K N V A N N S L K D F L
 CTGTCGTTTT TATTTTTGT AAATTCACTGC AAAAAAAATA ATGTATCATT GTCAATTGGT 840
 R D N K N K T F E H L F F L T D N D I P
 ATATTTCTGA TACTACTTTT GTTTTTGTT GGCAGGTATC TTTGGTTGAA ATGATAATCC 900
 I N R I S S K N K T P L Y R Q N F H Y D
 CAAGTTTTAT TAATTGATAA ATATTGTTA GTGTAATCAA TATCATTAAAC TGTAAACCT 960
 W T K N I S L Y K N T Y D I D N V T L G
 AACACATTCAAG CGAACCGCAT GCCAGTTTA GCGATGAGGT ATAACGCTGC ATACGATTGA 1020
 L C E A F R M
 <----|
 TGTTGTGATT TTTCTTACA AATTTTATC AAGCGTAAGT ATTCAATTGGT TTCAAGAAAT 1080
 TTATCTCTA TTACGCCCT TTATTTTG CTTAACCTT AGTGAATAAA CAAAAATTTT 1140
 TTTCTATATA TCCCTCGTGA ACAGCCATGG ATACGCAGGC TTTTACATGT ATGTTAAAC 1200
 GCTTTACTGT ATCTTGACACA TGCCTTGAC TATAATGATT TATGACTTGT TGATATTTAG 1260

TGGAAAGTAAT ATTGCAAAGT AATATATTC CTATTATATG TTTATACGAT ATTGATATT 1320
 CCCACCCGTT GTCGCGTTA CGGAAATACG CCATTGATAT ACTCCACATT AGCTAAAGAA 1380
 CAGGGTGTTC AAGGCTACCT TGATGGAAA GGCTCTCTA GAGATATTG TAAATGGTAT 1440
 GATATCTCAA GTCGCTCTGT TCTCCAAAAG TGGATAAAAC GGTATACTAG TGGTGAAGAC 1500
 TTGAAAGCCA CTAGTAGAGG ATATAGCCGT ATGAAACAAG GAAGGCAAGC CACATTTGAA 1560
 GAACGTGTAG AGATTGTTAA CTACACCATT GCCCATGGGA AAGACTATCA AGCAGCTATT 1620
 GAGAAAGTTG GTGTTCTCA CCAACAAATT TATTCTTGGG TGCGTAAGCT TGAGAAGAAT 1680
 GGCTCACAAG GTTGGTTGA TAGACGTGTG AAAGGGTTGG AGAGTAGGCC TGATTTAAC 1740
 GAGATTGAGC AACTTTAACT CAAGATTAAA CAATTGGAGG AACGTAATCG TCTCTTAGAA 1800
 ATCGAGGTTA GTTTACTAAA AAAGTTAGAA GACATCAAAC GAGGAAACAG ACGGTAAGAC 1860
 TAGGTAAGCA TTTAGCGGAG TTCCAAGTAA TCAAGAATTA TTACGATGAG GAATCTAATG 1920
 TGCCTATTCA GGCCCTATGC CAACTCTTGA AGGGGTCTCG TTCAAGGCTAT TACAAGTGGC 1980
 TCAATCGTCA AAAAACAGAT TTTGAGACAA AAAATACAAA GCTAATGGCT AAAATCAAGG 2040
 AACTTCGTAG ACTCTACAAT GGTATCTTAG GTTATCGCCG TATGACAACA TTTATTAATC 2100
 GTCAACTTGG GACAACCTAA ACAAGAAC GGATTCGTTG ATTGATGAAC ATTCTGGGA 2160
 TTAGTTCAAGT CATTGTCGT GTTAGCCATG CTTGTACAAA AGCTGGTGAC AGATTTACG 2220
 AAGAAAATAT TCTTAATCGT GAATTTACAG CCACAGCTCA TAACCAGAAA TGGTGCACAG 2280
 ATGTCACCTA TCTTCATAC GGTCTGGGAG CAAAGCTTA TCTCAGTGC ATAAAGACC 2340
 TGTATAACGG TTCTTATATC GCTTATGAGA TTAGTCACAA CAATGAAATC CACTTGTAT 2400
 GAAGACCATT AAAAAGGGC TAGAGCTAA TCCAGGAGCC ACACCTATCA TCCATAGCGA 2460
 TTGAGGTAGT CAATATACTT CCAAAGAATA CCGTTATATC ATACAACAAG CTGGTCTGAC 2520
 CTTATCCATG TCCCGGATTG GCAAATGTAT TGATAATGCA CCAACTGAA GTTCTTTGG 2580
 GTTTTCAAG ACTGAGTCTT ACCACCTAA GAAATACAAC TCTTATGATG AGTTGGTCAA 2640
 TGATGTGGCA CGTTATATCG AATTCTACAA CACACAACGT TATCAATCAA AATTAAACAA 2700
 CCTGACTCCT CTAGAATTCA GGAATCAGGT TGCATAACTT ATCTTTTATT ATTTGACTGT 2760
 CTACTTGACA GGGAGCCGTT CAGATTGCTT AACCTTCTA AATTCGCTAA AATAGCTACA 2820
 AGAAAACGAG CCATTTAATG CTTATTTCTT ATACTGTCTT GCCTCACGCT CTCCCTGGACC 2880
 AAAAATTGAG CGTGAGGCTT TTGTTTCAT TAAACGATGA TATTCGATA TTGATCAGTT 2940
 TGTTTCCGA GAGCCATCAA AGCTCGATA AGGTCGATAA TTCCAGGAAT AAAGGTAATA 3000
 CTAAAAATAA TATATAAAAA AACCTGGCCT ATTTTCCTG CGTAAATTT ATGCGCTCCA 3060
 ATGCCGCCCA AAAGAACGTT AATAAACAT AACTACTAT GTTAGCATAA GACTTTATTT 3120

TTACAACTGA ATTCATATA AATGGATTAG AGTAAGGGAT AAAAGAAATT AGCATAGCTC 3180
 TTTTAAAAAT AAAAAAATTA ATATAATATG GAAAAAATTT TATTCATAA ACGTTTCATA 3240
 AAAGGTATGT AATCTAGTAT TTAGGCAACA CTATTTGTC ACTGGTGTCT AGTAACCTAT 3300
 AGATTGATAA TTTTACTAGT AAACGTAATT CTCGCTTA AGAGTTAAAT GTCTATTTAT 3360
 TGTAAGCTAA ATTGGGAGGT GAACTTATGT AAAATTAGAT AGGTACTGTC AAGTACGGGA 3420
 TGATTATTGA AACAGCCAGT ATGCATCATA AAATCTGTAT TGCTTAATAA CTATTCCTT 3480
 AACAGACAT CAGTCATTG TTTATCATCG CTACCTAAG TCTAGTTTT TCAATAGAGC 3540
 ATTAGGTAGT TTTGATAAT AAAACTATAT AAACATGAGA ATTAGATTTC GTATTGCATT 3600
 CTTCATAATG AGTTATTGA GATTTCCCTT TGAATAAAATA GATACGAAAT TCAGTAACCT 3660
 CATATATAAA CGGCTCTATC ATTGAGATAG TTTGCAAAT GAAGAAATTT TTAATGGAAA 3720
 TAGTTTAAA AACATTAGTT GTAGGCGATG TAAAAATATT AATCCAGTGG ATGCAATAGT 3780
 TGCAGGAA AAATAGAGAG GAGTAATTAG GAAGTGATAA AAAATGCTAT AGCATATATT 3840
 ACCAGAAAAA AAAATAGAAC ACTTATTATA TTTGCTATTT TAACAATTGT TCTTCTTGC 3900
 TTGTATTCAAT GTTTAACAAAT AATGAAATCA AGTAATGAAA TAGAAAAGGC TTTATATGAA 3960
 M K S S N E I E K A L Y E
 |--->
 AGTTCTAATT CTTCAATATC AATTACAAAA AAAGATGGTA AATATTTAA TATTAATCAA 4020
 S S N S S I S I T K K D G K Y F N I N Q
 TTTAAGAATA TTGAAAAAAT AAAAGAGGTT GAAGAAAAAA TATTCATAA TGATGGATTA 4080
 F K N I E K I K E V E E K I F Q Y D G L
 GCAAAATTGA AAGATCTAA AGTAGTTAGT GGTGAGCAA GTATAATAG AGAAGATTTA 4140
 A K L K D L K V V S G E Q S I N R E D L
 TCTGACGAAT TTAAAAATGT TGTTCACTA GAAGCTACAA GTAATACTAA AAGAAATCTT 4200
 S D E F K N V V S L E A T S N T K R N L
 TTATTTAGTA GTGGAGTATT TAGTTTAAA GAAGGAAAAA ATATAGAAGA AAATGATAAG 4260
 L F S S G V F S F K E G K N I E E N D K
 AATTCAATTG TTGTTCATGA AGAATTGCT AAACAAAACA AACTAAATT GGGTGATGAA 4320
 N S I L V H E E F A K Q N K L K L G D E
 ATTGATCTTG AATTACTAGA TACGGAAAAA AGTGGAAAAA TAAAAAGTC TAAATTTAAA 4380
 I D L E L L D T E K S G K I K S H K F K
 ATTATAGGAA TCTTTCTGG TAAAAAACAG GAAACATATA CAGGATTATC ATCTGATTTT 4440
 I I G I F S G K K Q E T Y T G L S S D F
 AGCGAAAATA TGGTTTTGT AGATTATTCA ACTAGCCAAG AAATATAAA TAAATCAGAG 4500
 S E N M V F V D Y S T S Q E I L N K S E
 AATAATAGAA TTGCAAATAA AATTAAATG TATTCTGGTA GTTAAATC TACAGAGCTT 4560
 N N R I A N K I L M Y S G S L E S T E L
 GCCTTAAACA AATTGAAAGA CTTAAAATT GATAAGTCAA AGTATTCTAT TAAGAAAGAT 4620

A	L	N	K	L	K	D	F	K	I	D	K	S	K	Y	S	I	K	K	D
AATAAAGCAT TCGAAGAGTC TTTAGAGTC GTGAGTGGAA TAAAACATAT AATTAATAAATA 4680																			
N	K	A	F	E	E	S	L	E	S	V	S	G	I	K	H	I	I	K	I
ATGACTTATT CGATTATGTT AGGTGGAATA GTTGTCTTT CATTAATCTT GATTCTATGG 4740																			
M	T	Y	S	I	M	L	G	G	I	V	V	L	S	L	I	L	I	L	W
TTAAGAGAAA GAATTTATGA AATAGGTATA TTTTTATCTA TTGGAACAAC TAAGATACAA 4800																			
L	R	E	R	I	Y	E	I	G	I	F	L	S	I	G	T	T	K	I	Q
ATTATAAGGC AATTTATATT TGAGTTAATA TTCATATCAA TACCAAGTAT AATATCCTCC 4860																			
I	I	R	Q	F	I	F	E	L	I	F	I	S	I	P	S	I	I	S	S
TTATTTTTAG GGAATCTACT ATTAAAAAGTA ATTGTAGAAG GATTTATTAA CTCAGAGAAC 4920																			
L	F	L	G	N	L	L	L	K	V	I	V	E	G	F	I	N	S	E	N
TCAATGATT TCGGTGGAAG TTTAATAAT AAAAGCAGTT TTATGTTAAA CATAACAACA 4980																			
S	M	I	F	G	G	S	L	I	N	K	S	S	F	M	L	N	I	T	T
CTTGCAGAAA GTTATTTAAT ATTAATAAGT ATTATTGTTT TATCAGTTGT AATGGCTCT 5040																			
L	A	E	S	Y	L	I	L	I	S	I	I	V	L	S	V	V	M	A	S
TCATTAATAT TATTTAAGAA ACCACAAGAA ATATTATCAA AAATAAGTTA GGAGCAAATA 5100																			
S	L	I	L	F	K	K	P	Q	E	I	L	S	K	I	S	.			
ATGGATATAT TAGAAATAAA GAATGTAAT TACAGTTACG CAAATTCTAA AGAAAAAGTT 5160																			
M	D	I	L	E	I	K	N	V	N	Y	S	Y	A	N	S	K	E	K	V
----->																			
TTGTCAGGAG TAAATCAAAA ATTTGAACCTT GGAAAGTTTT ATGCGATAGT AGGGAAGTCA 5220																			
L	S	G	V	N	Q	K	F	E	L	G	K	F	Y	A	I	V	G	K	S
GGAACAGGAA AATCCACACT TCTTCCTTA CTTGCAGGAC TTGATAAAAGT TCAAACAGGA 5280																			
G	T	G	K	S	T	L	L	S	L	L	A	G	L	D	K	V	Q	T	G
AAAATCTTGT TTAAGAATGA AGATATAGAA AAGAAAGGAT ATAGTAATCA CAGAAAAAT 5340																			
K	I	L	F	K	N	E	D	I	E	K	K	G	Y	S	N	H	R	K	N
AATATATCTT TGGTATTCA AAATTATAAT TTAATAGATT ATTTATCGCC GATTGAAAAT 5400																			
N	I	S	L	V	F	Q	N	Y	N	L	I	D	Y	L	S	P	I	E	N
ATTAGACTAG TAAATAAAC T AGTAGATGAG AGTATCTTGT TCGAATTAGG TTTAGATAAA 5460																			
I	R	L	V	N	K	S	V	D	E	S	I	L	F	E	L	G	L	D	K
AAACAAATAA AAAGAAATGT TATGAAATTA TCTGGTGGTC AGCAACAAAG GGTAGCTATT 5520																			
K	Q	I	K	R	N	V	M	K	L	S	G	G	Q	Q	Q	R	V	A	I
GCTAGGGCAC TGGTATCAGA TGCCCCAATA ATACTAGCTG ATGAGCCTAC CGGTAACCTA 5580																			
A	R	A	L	V	S	D	A	P	I	I	L	A	D	E	P	T	G	N	L
GACAGTGTAA CTGCTGGAGA AATAATT (SEQ ID NO:27) 5607																			
D	S	V	T	A	G	E	I	I	.										

FIG. 5a

IQSLTEGQLR SDIPEFRAGD TVRVHAKVVE GTRERIQIFE GVVISRKGQG	50
ISEMYTVRKI SGGIGVERTF PIHTPRVDKI EVVRYGKVRR AKLYYLRALQ	100
GKAARIKEIR R (SEQ ID NO:28)	111

FIG. 5b

MRFAECLGLT VNDIDYTNKY LSINKTWDYH FNQRYLPTKN KSSIRNIPID	50
NDTLFFLHEF TKNKNDRLF D KLSNNAVNKT IRKITGREVR VHSLRHTFAS	100
YLISISQVLD HENLNITLEV YAHQLQEQQD RNDKLNQRNL GQNSSKPLFT	150
CNEYVPCRNR TSNYSLGGSC YIH (SEQ ID NO:29)	173

FIG. 5c

MKSSNEIEKA LYESSNSSIS ITKKDGKYFN INQFKNIEKI KEVEEKIFQY	50
DGLAKLKDLK VVSGEQSINR EDLSDEFKNV VSLEATSNTK RNLLFSSGVF	100
SFKEGKNIEE NDKNSILVHE EFAKQNLKL GDEIDLELLD TEKSGKIKSH	150
KFKIIGIFSG KKQETYTGLS SDFSENMVFV DYSTSQEILN KSENNRIANK	200
ILMYSGSLES TELALNKLKD FKIDKSKYSI KKDNKAFEEES LESVSGIKHI	250
IKIMTYSIML GGIWVLSLIL ILWLRERIYE IGIFLSIGTT KIQIIRQFIF	300
ELIFISIPSI ISSIFLGNLL LKVIVEGFIN SENSMIFGGS LINKSSFMLN	350
ITTLAESYLI LISIIVLSVV MASSLILFKK PQEILSKIS	389
(SEQ ID NO:30)	

FIG. 5d

MDILEIKNVN YSYANSKEKV LSGVNQKFEL GKFYAIVGKS GTGKSTLLSL	50
LAGLDKVQTG KILFKNEDIE KKGYSNHRKN NISL VFQNYN LIDYLSPIEN	100
IRLVNKSVDE SILFELGLDK KQIKRNMKL SGGQQQRVAI ARALVSDAPI	150
ILADEPTGNL DSVTAGEII (SEQ ID NO:31)	169

FIG. 5e

CATATGACAA TATTTTCAA AGTCTACATC ACTTACTCGC CTGTCGTGGA AAATCTGGCA	60
ATACATTAAT CGACCAATT A GTTGTGATG GTTACTTCA TGCAGATAAT CACTACCATT	120
TTTCAATGG GAAAGTCTCTG GCCACTTCA ATACTAACCA ATTGATTGCG GAAAGTTGTCT	180
ATGTTGAAAT ATCCTTAGAT ACTATGTCTA GTGGTGAACA TGATTTAGTA AAAGTTAACCA	240
TTATCAGACC CACTACCGAG CATACTATCC CCACGATGAT GACAGCTAGC CCCTATCATC	300
AAGGTATCAA TGATCTGCC GCAGACCAAA AAACATACCA AATGGAGGGT GCGCTAGCAG	360
TTAAACAGCC TAAACACATA CAAGTTGACA CAAAACCATT TAAAGAAGAA GTAAAACATC	420
CTTCAAAATT ACCCATCAGC CCTGCAACTG AAAGCTTCAC ACACATTGAC AGTTATAGTC	480
TCAATGACTA TTTTCTTCT CGTGGTTTG CTAATATATA CGTTTCAGGT GTGGGTACTG	540
CTGGCTCTAC GGGTTTCATG ACCAGTGGGG ATTACCAACA AATACAAAGC TTAAAGCAG	600
TCATTGATTG GTTAAATGGT AAGGTTACTG CATTCAACAAG TCATAAACAGA GATAAAACAAG	660
TCAAGGCTGA TTGGTCAAAC GGCCTTGTAG CAACCACAGG TAAATCTTAT CTCGGTACCA	720
TGTCAACTGG TTTAGCAACA ACTGGCGTTG AGGGGCTGAA AGTCATTATC GCTGAAGCCG	780
CAATCTCCAC ATGGTATGAT TATTATCGAG AAAATGGGCT TGTGTGAGT CCAGGCGGCT	840
ACCCCGGTGA AGATTTAGAC GTTTAACAG AATTAACATA CTCACGAAAC CTCTTAGCTG	900
GTGATTACAT CAAAAACAAAC GATTGCTATC AAGCATTGTT AAATGAACAA TCAAAAGCAA	960
TTGACCGTCA AAGTGGGGAT TACAACCAAT ACTGGCATGA CCGTAATTAC CTAACTCACG	1020
TCAATAATGT CAAAAGTCGA GTAGTTTACA CTCATGGACT ACAGGATTGG AATGTTAACG	1080
CAAGACATGT CTACAAAGTT TTCAATGCAT TGCCTCAAAC CATAAAAAA CACCTTTTT	1140
TACATCAAGG TCAACATGTG TATATGCATA ATTGGCAGTC GATTGATTTC CGTGAAAGCA	1200
TGAATGCCTT ACTAACGCAA GAACTACTTG GCATTGACAA TCATTCCAA TTAGAAGAGG	1260
TCATTTGGCA AGATAAACT ACTGAGCAA CTTGGCAAGT TTTAGATGCT TTGGGAGGAA	1320
ACCATCAAGA GCAAATTGGT TTAGGTGATA GTAAAAAAACT TATTGATAAC CATTATGACA	1380
AAGAAGCCTT TGATACTTAT TGAAAGACT TCAATGTGTT CAAAATGAT CTTTCAGG	1440
GAAATAATAA AACCAATCAA ATCACTATTA ATCTTCCTCT AAAGAAAAAT TATCTCCTGA	1500
ATGGACAGTG CAAACTCCAT CTACGTGTTA AAACTAGTGA CAAAAGGCC ATTTTATCAG	1560
CCCAAATCTT AGACTATGGT CCTAAAAAAC GATTCAAAGA TACACCAACC ATCAAATTCT	1620
TAAACAGCCT TGATAATGGT AAAAATTTG CCAGAGAACG TTTACGTGAA CTCCCGTTA	1680
CTAAAGATCA TTATCGTGTCA ATCAGTAAAG GTGTCTTGAA CCTTCAAAAT CGTACAGACT	1740
TACTTACAAT TGAGGCTATC GAGCCAGAAC AATGGTTGA TATCGAGTTT AGCCTCCAA	1800
CAAGTATATA TCAATTGAGT AAAGGTGATA ATCTAAGGAT TATCCTTTAT ACAACTGATT	1860
TTGAACATAC CATTGAGAT AATGCTAGTT ACTCTATAAC AGTAGATTTG AGTCAATCTT	1920
ATTTAACTAT CCCAACTAAT CAAGGAAATT AACTTATGAA ACTTCTTACT AAAGAACGGT	1980
TTGATGATTC TCAACACTTT TGGTACCAAGA TCAATTATT ACAAGAGAGT AACTTCGGAG	2040
CAGTTTTGA CCATGATAAT AAAAACATTC CACAGGTTGT TGCAACTATT GTTGATGATT	2100
TACAAGGTTTC CGGAAGTTCG AATCATTCTT GGTATTTGG CAATACTACT GATACTTCCA	2160
TCCTTATGAT TGCTCATTTA AATCGAAAT TCTATATTCA GGTTAATTAA AAGGACTTTG	2220
ACTTTGCACT CAATTAAATA GCTATAAATA ATTGGAAGAG TCTCCTCCAA ACTCAACTTG	2280
AAGCTCTAAA CGATACCCCTA GCAATATTTC AATAAATAAG GTAGAATGGG GTGACAAAGC	2340
AACCGCAGGG AGACTGATTA ATGTCATCTT ATTGGAATAA CTATCCTGAA CTTAAAAAAA	2400

ATATTGATGA AACCAATCAA CTAATTCAAG AAAGAATACA GGTAGAAAT AAAGATATTG 2460
 AAGCGGCCT AAGCCAACTC ACAGCTGCGG GAGGAAAACA GCTCAGACCA GCATTCTTT 2520
 ACCTTTTTC TCAACTTGGT AATAAGGAGA ATCAAGATAC TCAGCAACTA AAGAAAATCG 2580
 CTGCTTCTT AGAAATCCTT CACGTTGCTA CATTAAATCCA TGATGATGTC ATTGATGACT 2640
 CACCACTAAG ACGTGAAAT ATGACCATTG AAAGCAAGTT TGGCAAAGAC ATCGCAGTT 2700
 ATACTGGGGA TTTACTTTTC ACAGTCTTT TCGATCTTAT TTTAGAATCT ATGACTGATA 2760
 CACCATTTAT GAGGATTAAT GCAAAATCTA TCGTAAAAT TCTCATGGG GAATTGGACC 2820
 AGATGCACCT TCGTTACAAT CAACAACAAG GTATCCATCA CTATTTACGT GCGATTTCA 2880
 GTAAGACAGC CGAACTCTT AAATTAGCTA GCAAAAGAAGG AGCTTACTTT GGTGGTGCAG 2940
 AGAAGGGAGGT TGTTCGTCTA GCAGGCCATA TCGGCTTTAA CATTGGTATG ACATTCCAAA 3000
 TTTTGGATGA TATCCTGGAT TATACTGCAG ATAAAAAAAC ATTTAATAAG CCTGTCTTAG 3060
 AGGATTTAAC ACAAGGCCTT TACAGCCTTC CTCTACTTCT TGCCATTGAA GAAAATCCTG 3120
 ATATTTCAA ACCTATTTA GATAAAAAAA CAGATATGGC TACTGAAGAC ATGGAAAAAA 3180
 TTGCTTATCT CGTCGTTCC CATAGAGGTG TTGACAAAGC TCGCCATCTA GCTCGTAAAT 3240
 TTACTGAGAA AGCTATTAGT GACATAAATA AGCTACCCCA GAACTCTGCA AAAAAACAGT 3300
 TGCTACAATT AACTAATTAC CTTTTAAAAC GCAAAATTAA AATAAAAAA AAACATTCCA 3360
 CAATGCTAGA AAAGCAGTTA GGGATGTTT TTTTATTATC ATTTATTTAT CGCACCTATC 3420
 AATCATCATA GATCACCACAT ATCAGCGGCT TTCAGCTGAC GGTAACGTTG ACTACTTGA 3480
 GACAATTCTT GAGGAGAACC TTCCAACCT AATTGCCAT TTTCTATAAA TAAGATACGA 3540
 TCAGCATGTT CAATACCTT TAAGTGTGTA GTAAATCCAAA CTAAGGTCTT ACCTTCCAAT 3600
 TCTTCATAA ATACCCCTAG TAAGGCTTGT TCAGTAATAG GATCAAGTCC AACAGTTGGC 3660
 TCATCTAAGA TAACAATTGG GACATCTTT AGTAAGATTC TAGCCAAAGC AATTCTATGC 3720
 CTTTCGCCAC CTGAAAACCT AAGTCCAGCT TCATCAACCA TTGTATAGAG ACCATCTGAT 3780
 AAATCAGTGA CCATCTCTT CAATCCAACCT CGTTCAAGAA CTTTCCATAC ATCTTCTTCA 3840
 CTAGCATCTT GGTTCCAAT GCGAATGTTA TTTAGCAGGG TTGTATTTAA AAGGTAGGGC 3900
 GCTTGTGTA TCACTCCAAT ATAGTTGAA ATGCAATCAC CAACTATTGA AACATCAGCA 3960
 CCGCCTAGGG TAATCTCCC TTGACTTGCT TTCAAGTCGC CACGAAGTAG ACTAGCTAAG 4020
 GTACTCTTGC CAGAACCACT CCGCCCTAAA ATAGCAATT TTTCTCCTTC TTTAATATCC 4080
 AAATCTAAAT GATGCAAACAC CCATTTCTCT TGTGGCTTAT ACTGGAAACT TAAATTCTTG 4140
 ACGGAAAAT CATATGGCTT ATTAGGCAAT T (SEQ ID NO:32) 4171

FIG. 6a

YDNIFQSLHH LLACRGKSGN TLIDQLVADG LLHADNHYHF FNGKSLATFN 50
 TNQLIREVVY VEISLDTMSS GEHDLVKVNI IRPTTEHTIP TMMTASPYHQ 100
 GINDPAADQK TYQMEGALAV KQPKHIQVDT KPFKEEVKHP SKLPISPATE 150
 SFTHIDSYSL NDYFLSRGFA NIYVSGVGTA GSTGFMTSGD YQQIQSFKAV 200
 IDWLNGKVTA FTSHKRDQV KADWSNGLVA TTGKSYLGTM STGLATTGVE 250
 GLKVIIAEAA ISTWYDYYRE NGLVCSPGGY PGEDLDVLTE LTYSRNLLAG 300
 DYIKNNDCYQ ALLNEQSKAI DRQSGDYNQY WHDRNYLTHV NNVKSRRVYVT 350
 HGLQDWNVKP RHVYKVFNAL PQTIKKHLFL HQGQHVMHN WQSIDFRESM 400
 NALLSQELLG IDNHFQLEEV IWQDNTTEQT WQVLDAGGN HQEQIGLGDS 450
 KKLIDNHYDK EAFDTYCKDF NVFKNDLFKG NNKTNQITIN LPLKKNYLLN 500
 GQCKLHLRVK TSDKKAILS QILDYGPKKR FKDTPTIKFL NSLDNGKNFA 550
 REALRELPFT KDHYRVISKG VLNLQNRTDL LTIEAIEPEQ WFDIEFSLQP 600
 SIYQLSKGDN LRIILYTTDF EHTIRDNASY SITVDSLQSY LTIPTNQGN 649
 (SEQ ID NO:33)

FIG. 6b

MKLLTKERFD DSQHFWYQIN LLQESNFGAV FDHDNKNIPQ VVATIVDDLQ 50
 GSGSSNHFWY FGNTTDTTSIL MIAHLNRKFY IQVNLKDFDF ALNLIAINNW 100
 KSLLQTQLEA LNDTLAIFQ (SEQ ID NO:34) 119

FIG. 6c

MSSYWNNYPE LKKNIDETNQ LIQERIQVRN KDIEAALSQL TAAGGKQLRP 50
 AFFYLFSQLG NKENQDTQQL KKIAASLEIL HVATLIHDDV IDDSPLRRGN 100
 MTIQSKFGKD IAVYTGDLLF TVFFDLILES MTDTPFMRIN AKSMRKILMG 150
 ELDQMHLRYN QQQGIHHYLR AISGKTAELF KLASKEGAYF GGAEKEVVRL 200
 AGHIGFNIGM TFQILDDILD YTADKKTFNK PVLEDLTQGV YSLPLLLAIE 250
 ENPDIFKPIL DKKTDMATED MEKIAYLVVS HRGVDKARHL ARKFTEKAIS 300
 DINKLPQNSA KKQLLQLTNY LLKRKI (SEQ ID NO:35) 326

FIG. 6d

LPNKPYDFSV KNLSFQYKPQ EKWVLHHLDL DIKEGEKIAI LGRSGSGKST 50
LASLLRGDLK ASQGKITLGG ADVSIVGDCI SNYIGVIQQA PYLFNTTLLN 100
NIRIGNQDAS EEDVWKVLER VGLKEMVTDL SDGLYTMVDE AGLRFSGGER 150
HRIALARILL KDVPIVILDE PTVGLDPITE QALLRVFMKE LEGKTLVWIT 200
HHLKGIEHAD RILFIENGQL ELEGSPQELS QSSQRYRQLK AADDGDL 247
(SEQ ID NO:36)

FIG. 6e

AATTCTATTT GGAGGTTTT CTTGAATAAA TGGTTAGTTA AGGCAAGTTC CTTAGTTGTT	60
TTAGGTGGTA TGGTTTTATC TGCGGGTTC CGAGTTTAG CGGATACTTA TGTCCGTCCA	120
ATTGATAATG GTAGAATTAC AACAGGTTTC AATGGTTATC CTGGACATTG TGGGGTGGAT	180
TATGCTGTTG CGACTGGAAC GATTATTAGG GCAGTGGCAG ATGGTACTGT GAAATTGCA	240
GGAGCTGGAG CCAACTTTTC TTGGATGACA GACTTAGCAG GAAATTGTGT CATGATTCAA	300
CATGCGGATG GAATGCATAG TGTTACGCT CATATGTCAC GTGTGGTGGC TAGGACTGGG	360
GAAAAAGTCA AACAGGAGA TATCATCGGT TACGTAGGAG CAACTGGTAT GGCGACGGGA	420
CCTCACCTTC ATTTGAATT TTTACCAAGCT AACCTTAATT TTCAAAATGG TTTCCATGGA	480
CGTATCAATC CAACGTCACT AATTGCTAAC GTTGCACCT TTAGTGGAAA AACGCAAGCA	540
TCAGCTCCAA GCATTAAGCC ATTACAATCA GCTCCTGTAC AGAATCAATC TAGTAAATTA	600
AAAGTGTATC GAGTAGATGA ATTACAAAAG GTTAATGGTG TTTGGTTAGT CAAAAATAAC	660
ACCCCTAACCG CGACTGGGTT TGATTGGAAC GATAATGGTA TACCAGCATIC AGAAATTGAT	720
GAGGTTGATG CTAATGGTAA TTTGACAGCT GACCAGGTTTC TTCAAAAGG TGGTTACTTT	780
ATCTTTAATC CTAAAACCTCT TAAGACTGTA GAAAACCCA TCCAAGGAAC AGCTGGTTA	840
ACTTGGGCTA AGACACGCTT TGCTAATGGT AGTCAGTTT GGCTTCGCGT TGACAACAGT	900
CAAGAACTGC TTTACAAATAA GTTGAGGTA TTGATTCAATT GTTTAAATG ACAGTTTGT	960
TACTAACTAA GTACAATTTCTTAAACCGT CTGAAAATAA TTTTATAGTC CAGTAAAGTG	1020
TGATATTATA GTCTCGGACT AATAAAAAGG AAATAGGAAT TGAAGCAATG AAAATGAATA	1080
AAAAGGTACT ATTGACATCG ACAATGGCAG CTTCGCTATT ATCAGTCGCA AGTGTCAAG	1140
CACAAGAAC AGATACGACG TGGACAGCAC GTACTGTTTC AGAGGTAAAG GCTGATTG	1200
TAAGCAAGA CAATAAATCA TCATATACTG TGAAATATGG TGATACACTA AGCGTTATTT	1260
CAGAAGCAAT GTCAATTGAT ATGAATGTCT TAGCAAAAT TAATAACATT GCAGATATCA	1320
ATCTTATTAA TCCGTGAGACA ACACTGACAG TAACTTACGA TCAGAAGAGT CATACTGCCA	1380
CTTCAATGAA AATAGAAAACA CCAGCAACAA ATGCTGCTGG TCAAACAACA GCTACTGTGG	1440
ATTTGAAAAC CAATCAAGTT TCTGTTGCAG ACCAAAAAGT TTCTCTCAAT ACAATTTCGG	1500
AAGGTATGAC ACCAGAAGCA GCAACAACGA TTGTTTCGCC AATGAAGACA TATTCTTCTG	1560
CGCCAGCTTT GAAATCAAAA GAAGTATTAG CACAAGAGCA AGCTGTTAGT CAAGCAGCAG	1620
CTAATGAACA GGTATCAACA GCTCCTGTGA AGTCGATTAC TTCAGAAGTT CCAGCAGCTA	1680
AAGAGGAAGT TAAACCAACT CAGACGTCAG TCAGTCAGTC AACAAACAGTA TCACCAGCTT	1740
CTGTTGCCGC TGAAACACCA GCTCCAGTAG CTAAAGTAGC ACCGGTAAGA ACTGTAGCAG	1800
CCCCTAGAGT GGCAAGTGTT AAAGTAGTCA CTCCTAAAGT AGAAACTGGT GCATCACCAG	1860
AGCATGTATC AGCTCCAGCA GTTCCGTGTA CTACGACTTC AACAGCTACA GACAGTAAGT	1920
TACAAGCGAC TGAAAGTTAAG AGCGTTCCGG TAGCACAAAA AGCTCCAACA GCAACACCGG	1980
TAGCACAAACC AGCTTCAACA ACAAAATGCAG TAGCTGCACA TCCCTAAAAT GCAGGGCTCC	2040
AACCTCATGT TGCAGCTTAT AAAGAAAAAG TAGCGTCAAC TTATGGAGTT AATGAATTCA	2100
GTACATACCG TGCAGGTGAT CCAGGTGATC ATGGTAAAGG TTTAGCAGTC GACTTTATTG	2160
TAGGTAAAAA CCAAGCACTT GGTAAATGAAG TTGCACAGTA CTCTACACAA AATATGGCAG	2220
CAAATAACAT TTCATATGTT ATCTGGCAAC AAAAGTTTA CTCAAATACA AATAGTATTT	2280
ATGGACCTGC TAATACTTGG AATGCAATGC CAGATCGTGG TGGCCTTACT GCCAACATT	2340
ATGACCATGT TCACGTATCA TTTAACAAAT AATATAAAA AGGAAGCTAT TTGGCTTCTT	2400

TTTTATATGC CTTGAATAGA CTTCAAGGT TCTTATCTAA TTTTTATTAA ATTGAGGAGA 2460
 TTAAGCTATA AGTCTGAAAC TACTTCACG TTAACCGTGA CTAATCAAA ACGTTAAAAC 2520
 TAAAATCTAA GTCTGTAAAG ATTATTGAAA ACGTTAAA AACAGATATA ATAAGGTTG 2580
 TAGATATCTA AAATTAAGGAA AGATAAGGAA GTGAGAATAT GCCACATCTA AGTAAAGAAG 2640
 CTTTAAAGA GCAAATAAAAA AATGGCATT TTGTGTCAATG TCAAGCTTG CCTGGGGAGC 2700
 CTCTTATAC TGAAAGTGGG GGTGTTATGC CTCTTTAGC TTTGGCAGCT CAAGAAGCAG 2760
 GAGCGGTTGG TATAAGAGCC AATAGTGTCC GCGACATTAA GGAAATTCAA GAAGTTACTA 2820
 ATTTACCTAT CATCGGCATT ATAAACGTG AATATCCTCC ACAAGAACCA TTTATCACTG 2880
 CTACGATGAC AGAGGTGGAT CAATTAGCTA GTTTAGATAT TGCAAGTAATA GCCTTAGATT 2940
 GTACACTTAG AGAGCGTCAT GATGGTTGA GTGTAGCTGA GTTTATTCAA AAGATAAAAG 3000
 GGAAATATCC TGAACAGTTG CTAATGGCTG ATATAAGTAC TTTGAAGAA GGTAAGGATG 3060
 CTTTGAAGC AGGAGTTGAT TTTGTGGTA CAACTCTATC TGGATACACA GATTACAGCC 3120
 GCCAAGAAGA AGGACCGGAT ATAGAACTCC TTAATAAGCT TTGTCAAGCC GGTATAGATG 3180
 TGATTGCGGA AGGTAAAATT CATACTCCTA AGCAAGCTAA TGAAATTAAT CATATAGGTG 3240
 TTGCAGGAAT TGTAGTTGGT GGTGCTATCA CTAGACCAAA AGAAATAGCG GAGCGTTCA 3300
 TCTCAGGACT TAGTTAAAAG TGTACTCAA AAATCAAAAT CAAATAAAAA AAGGGAAATA 3360
 GTTATGAGTA TCAAAAAAAG TGTGATTGGT TTTGCCTCG GAGCTGCAGC ATTATCAATG 3420
 TTTGCTTGTG TAGACAGTAG TCAATCTGTT ATGGCTGCCG AGAAGGATAA AGTCGAAATT 3480
 (SEQ ID NO:37)

FIG. 7a

NSIWRFFLNK WLVKASSLVV LGGMVL SAGS RVLADTYVRP IDNGRITTGF 50
 NGYPGHCGVD YAVPTGTIIR AVADGTVKFA GAGANFSWMT DLAGNCVMIQ 100
 HADGMHSGYA HMSRVVARTG EKVKGDIIG YVGATGMATG PHLHFEFLPA 150
 NPNFQNGFHG RINPTSLIAN VATFSGKTQA SAPSIKPLQS APVQNQSSKL 200
 KVYRVDELQK VNGVWLKNN TLTPTGFDWN DNGIPASEID EVDANGNLTA 250
 DQVLQKGGYF IFNPKTLKTV EKPIQGTAGL TWAKTRFANG SSVWLRVDNS 300
 QELLYK (SEQ ID NO:38) 306

FIG. 7b

MKMNKKVLLT	STMAASLLSV	ASVQAQETDT	TWTARTVSEV	KADLVKQDNK	50
SSYTVKYGDT	LSVISEAMSI	DMNVLAKINN	IADINLIYPE	TTLTVTYDQK	100
SHTATSMKIE	TPATNAAGQT	TATVDLKTNQ	VSVADQKVSL	NTISEGMTPE	150
AATTIVSPMK	TYSSAPALKS	KEVLAQEQAQ	SQAAANEQVS	TAPVKSITSE	200
VPAAKEEVKP	TQTSVSQSTT	VSPASVAAET	PAPVAKVAPV	RTVAAPRVAS	250
VKVVTPKVET	GASPEHVSAP	AVPVTTSTA	TDSKLQATEV	KSVPVAQKAP	300
TATPVAQPAS	TTNAVAAHPE	NAGLQPHVAA	YKEKVASTYQ	VNEFSTYRAG	350
DPGDHGKGLA	VDFIVGKNQA	LGNEVAQYST	QNMAANNISY	VIWQQKFYSN	400
TNSIYGPANT	WNAMPDRGGV	TANHYDHVHV	SFNK	(SEQ ID NO:39)	434

FIG. 7c

MPHLSKEAFK	KQIKNGIIVS	CQALPGEPLY	TESGGVMPLL	ALAAQEAGAV	50
GIRANSVRDI	KEIQEVTNLP	IIGIIKREYP	PQEPMFITATM	TEVDQLASLD	100
IAVIALDCTL	RERHDGLSVA	EFIQKIKGKY	PEQLLMADIS	TFEEGKNAFE	150
AGVDFVGTTL	SGYTDYXRQE	EGPDIELLNK	LCQAGIDVIA	EGKIHTPKQA	200
NEINHIGVAG	IVVGGAITRP	KEIAERFISG	LS	(SEQ ID NO:40)	232

FIG. 7d

MSIKKSVIGF	CLGAAALSMF	ACVDSSQSVM	AAEKDKVEI	39
(SEQ ID NO:41)				

FIG. 7e

ATGAAAATGA	ATAAAAAAGGT	ACTATTGACA	TCGACAATGG	CAGCTTCGCT	50
ATTATCAGTC	GCAAGTGTTC	AAGCACAAGA	AACAGATAACG	ACGTGGACAG	100
CACGTACTGT	TTCAGAGGTA	AAGGCTGATT	TGGTAAAGCA	AGACAATAAA	150
TCATCATATA	CTGTGAAATA	TGGTGATACA	CTAACCGTTA	TTTCAGAAGC	200
AATGTCAATT	GATATGAATG	TCTTAGCAAA	AATTAATAAC	ATTGCAGATA	250
TCAATCTTAT	TTATCCTGAG	ACAACACTGA	CAGTAACCTTA	CGATCAGAAG	300
AGTCATACTG	CCACCTCAAT	GAAAATAGAA	ACACCCAGCAA	CAAATGCTGC	350
TGGTCAAACAA	ACAGCTACTG	TGGATTGAA	AACCAATCAA	GTTTCTGTTG	400
CAGACCAAAA	AGTTTCTCTC	AATACAATT	CGGAAGGTAT	GACACCAGAA	450
GCAGCAACAA	CGATTGTTTC	GCCAATGAAG	ACATATTCTT	CTGCGCCAGC	500
TTTGAAATCA	AAAGAAGTAT	TAGCACAAGA	GCAAGCTGTT	AGTCAAGCAG	550
CAGCTAATGA	ACAGGTATCA	ACAGCTCCTG	TGAAGTCGAT	TACTTCAGAA	600
GTTCCAGCAG	CTAAAGAGGA	AGTTAAACCA	ACTCAGACGT	CAGTCAGTCA	650
GTCAACAACA	GTATCACCGAG	CTTCTGTTGC	CGCTGAAACA	CCAGCTCCAG	700
TAGCTAAAGT	AGCACCGGT	AGAACTGTAG	CAGCCCCTAG	AGTGGCAAGT	750
GTTAAAGTAG	TCACTCCTAA	AGTAGAAACT	GGTGCATCAC	CAGAGCATGT	800
ATCAGCTCCA	GCAGTTCTG	TGACTACGAC	TTCAACAGCT	ACAGACAGTA	850
AGTTACAAGC	GACTGAAGTT	AAGAGCGTTC	CGGTAGCACA	AAAAGCTCCA	900
ACAGCAACAC	CGGTAGCACA	ACCAGCTTC	ACAACAAATG	CAGTAGCTGC	950
ACATCCTGAA	AATGCAGGGC	TCCAACCTCA	TGTTGCAGCT	TATAAAGAAA	1000
AAGTAGCGTC	AACTTATGGA	GTAAATGAAT	TCAGTACATA	CCGTGCAGGT	1050
GATCCAGGTG	ATCATGGTAA	AGGTTTAGCA	GTCGACTTTA	TTGTAGGTAA	1100
AAACCAAGCA	CTTGGTAATG	AAGTTGCACA	GTACTCTACA	CAAATATGG	1150
CAGCAAATAA	CATTTCATAT	GTTATCTGGC	AACAAAAGTT	TTACTCAAAT	1200
ACAAATAGTA	TTTATGGACC	TGCTAATACT	TGGAATGCAA	TGCCAGATCG	1250
TGGTGGCGTT	ACTGCCAAC	ATTATGACCA	TGTTCACGTA	TCATTTAACAA	1300
AATAA					1305

(SEQ ID NO:42)

FIG. 8

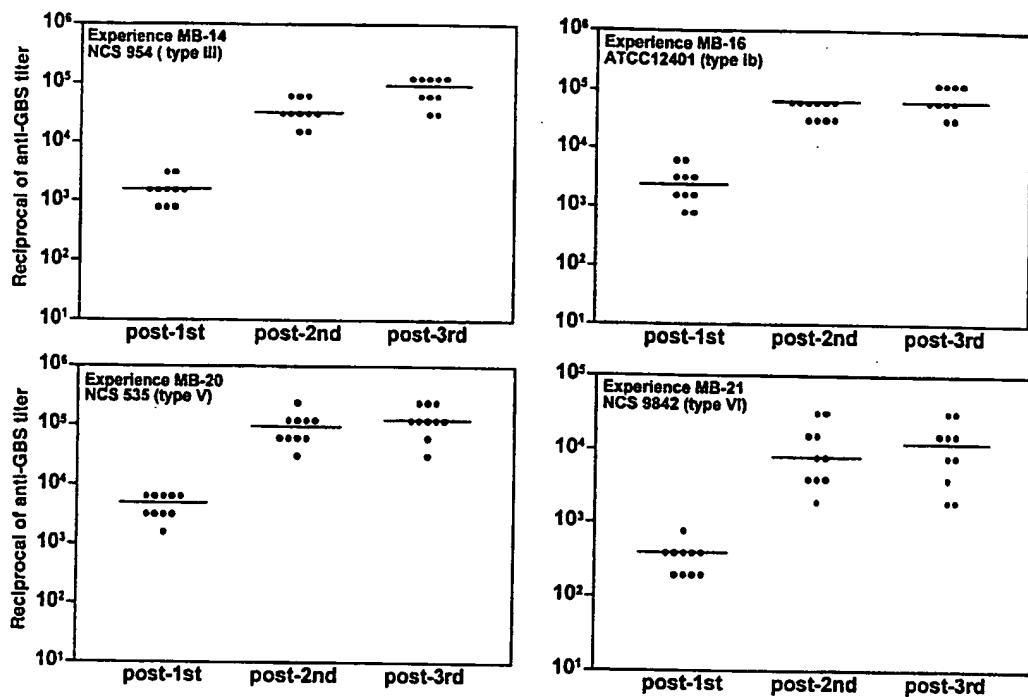
CAAGAAACAG	ATACGACGTG	GACAGCACGT	ACTGTTTCAG	AGGTAAAGGC	50
TGATTTGGTA	AAGCAAGACA	ATAAACATC	ATATACTGTG	AAATATGGTG	100
ATACACTAAG	CGTTATTTCA	GAAGCAATGT	CAATTGATAT	GAATGTCTTA	150
GCAAAAATTA	ATAACATTGC	AGATATCAAT	CTTATTATAC	CTGAGACAAC	200
ACTGACAGTA	ACTTACGATC	AGAAGAGTCA	TAATGCCACT	TCAATGAAAA	250
TAGAAACACC	AGCAACAAAT	GCTGCTGGTC	AAACAACAGC	TACTGTGGAT	300
TTGAAAACCA	ATCAAGTTTC	TGTTGCAGAC	CAAAAGTTT	CTCTCAATAC	350
AATTCGGAA	GGTATGACAC	CAGAAGCAGC	AACAACGATT	GTTTCGCCAA	400
TGAAGACATA	TTCTTCTGCG	CCAGCTTGA	AATCAAAGA	AGTATTAGCA	450
CAAGAGCAAG	CTGTTAGTCA	AGCAGCAGCT	AATGAACAGG	TATCAACAGC	500
TCCTGTGAAG	TCGATTACTT	CAGAAGTTCC	AGCAGCTAAA	GAGGAAGTTA	550
AACCAACTCA	GACGTCAGTC	AGTCAGTCAA	CAACAGTATC	ACCAGCTTCT	600
GTTGCCGCTG	AAACACCAGC	TCCAGTAGCT	AAAGTAGCAC	CGGTAAAGAAC	650
TGTAGCAGCC	CCTAGAGTGG	CAAGTGTAA	AGTAGTCACT	CCTAAAGTAG	700
AAACTGGTGC	ATCACCAGAG	CATGTATCAG	CTCCAGCAGT	TCCTGTGACT	750
ACGACTTCAA	CAGCTACAGA	CAGTAAGTTA	CAAGCGACTG	AAGTTAAGAG	800
CGTTCGGTA	GCACAAAAAG	CTCCAACAGC	AACACCGGTA	GCACAACCAAG	850
CTTCACAAAC	AAATGCAGTA	GCTGCACATC	CTGAAAATGC	AGGGCTCCAA	900
CCTCATGTTG	CAGCTTATAA	AGAAAAAGTA	GCGTCAACTT	ATGGAGTTAA	950
TGAATTCACT	ACATACCGTG	CAGGTGATCC	AGGTGATCAT	GGTAAAGGTT	1000
TAGCAGTCGA	CTTTATTGTA	GGTAAAAACC	AAGCACTTGG	TAATGAAGTT	1050
GCACAGTACT	CTACACAAAA	TATGGCAGCA	AATAACATT	CATATGTTAT	1100
CTGGCAACAA	AAGTTTTACT	CAAATACAAA	TAGTATTAT	GGACCTGCTA	1150
ATACTTGAA	TGCAATGCCA	GATCGTGGTG	GCGTTACTGC	CAACCATTAT	1200
GACCATGTTCA	ACGTATCATT	TAACAAATAA	(SEQ ID NO:43)		1230

FIG. 9

QETDTTW TAR	TVSEVKADLV	KQDNKSSYTV	KYGD TLSVIS	EAMSID MNVL	50
AKINNIADIN	LIYPETTLTV	TYDQKSHTAT	SMKIE TPATN	AAGQTTATVVD	100
LKTNQVSVAD	QKVSLNTISE	GMTPEAATTI	VSPM KTYSSA	PALKSKEVLA	150
QEQA VSQAAA	NEQVSTAPVK	SITSEVPAAK	EEVKPTQTSV	SQSTTVSPAS	200
VAAE TPAPVA	KVAPVRTVAA	PRVASVKVVT	PKVETGASPE	HVSAPAVPVT	250
TTSTATDSKL	QATEVKSVPV	AQKAPTATPV	AQPA STTNAV	AAHPENAGLQ	300
PHVAAYKEKV	ASTYGVNEFS	TYRAGDPGDH	GKGLAVDFIV	GKNQALGNEV	350
AQYSTQNMAA	NNISYVIWQQ	KFYSNTNSIY	GPANTWNAMP	DRGGVTANHY	400
DHVHVSFNK	(SEQ ID NO:44)				409

FIG. 9a

Fig. 10



SEQUENCE LISTING

<110> BioChem Vaccins
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HAMEL, Josée
CHARLEBOIS, Isabelle
BOYER, Martine

<120> NOVEL GROUP B STREPTOCOCCUS ANTIGENS

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tac ttt att atc cta ttg aca gca tgg act ttg atg gca tta gta act Tyr Phe Ile Ile Leu Leu Thr Ala Trp Thr Leu Met Ala Leu Val Thr 50 55 60	191
gct tta gtg gga tgg gat aat agg tat ggt tcc ttc ttg tcg tta tta Ala Leu Val Gly Trp Asp Asn Arg Tyr Gly Ser Phe Leu Ser Leu Leu 65 70 75	239
ata tta tta ttc cag ctt ggt tca agc gca gga act tac cca ata gaa Ile Leu Leu Phe Gln Leu Gly Ser Ser Ala Gly Thr Tyr Pro Ile Glu 80 85 90 95	287
ttg agt cct aag ttc ttt caa aca att caa cca ttt tta ccg atg act Leu Ser Pro Lys Phe Phe Gln Thr Ile Gln Pro Phe Leu Pro Met Thr 100 105 110	335
tac tct gtt tca gga tta aga gag acc atc tcg ttg acg gga gac gtt Tyr Ser Val Ser Gly Leu Arg Glu Thr Ile Ser Leu Thr Gly Asp Val 115 120 125	383
aac cat caa tgg aga atg cta gta atc ttt tta gta tca tcg atg ata Asn His Gln Trp Arg Met Leu Val Ile Phe Leu Val Ser Ser Met Ile 130 135 140	431
ctt gct ctt ctt att tat cgt aaa caa gaa gat taatagaaaag tatctagtga Leu Ala Leu Leu Ile Tyr Arg Lys Gln Glu Asp 145 150	484
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caa ccg att gtt ttt tta cat ggc aat agc tta agt agt cgc tat ttt Gln Pro Ile Val Phe Leu His Gly Asn Ser Leu Ser Ser Arg Tyr Phe 295 300 305 310	1137
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ttc agg caa ata gca gtt gac tta aag gat atc tta gtt cat tta gag Phe Arg Gln Ile Ala Val Asp Leu Lys Asp Ile Leu Val His Leu Glu 345 350 355	1281
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aat tca ggg aac ctg act att cat ggt cag cga tgg tgg gat att ctt Asn Ser Gly Asn Leu Thr Ile His Gly Gln Arg Trp Trp Asp Ile Leu 395 400 405	1425
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ccg tat atg agg caa aaa gct caa gtt att tcg ctt atg ttg gag gat 1521
 Pro Tyr Met Arg Gln Lys Ala Gln Val Ile Ser Leu Met Leu Glu Asp
 425 430 435

 ttg aag att agt cca gct gat tta cag cat gtg tca act cct gta atg 1569
 Leu Lys Ile Ser Pro Ala Asp Leu Gln His Val Ser Thr Pro Val Met
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 Val Leu Val Gly Asn Lys Asp Ile Ile Lys Leu Asn His Ser Lys Lys
 455 460 465 470

 ctt gct tct tat ttt cca agg ggg gag ttt tat tct tta gtt ggc ttt 1665
 Leu Ala Ser Tyr Phe Pro Arg Gly Glu Phe Tyr Ser Leu Val Gly Phe
 475 480 485

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 Gly His His Ile Ile Lys Gln Asp Ser His Val Phe Asn Ile Ile Ala
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 Lys Lys Phe Ile Asn Asp Thr Leu Lys Gly Glu Ile Val Glu Lys Ala
 505 510 515

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 Leu Ile Ile Leu Val Gly Val Ile Ala Val Leu Pro Thr Thr Gly Tyr
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 Asp Phe Val Leu Asn Gly Leu Leu Arg Thr Asp Lys Ser Lys Arg Tyr
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 Ile Leu Gln Thr Ser Trp Cys Ile Asn Thr Phe Asn Asn Leu Ser Gly
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 ttc ggt ggc tta atc gat att ggg ttg cgc atg gct ttt tat ggt aaa 2110
 Phe Gly Gly Leu Ile Asp Ile Gly Leu Arg Met Ala Phe Tyr Gly Lys
 595 600 605

 aaa ggt caa gag aag agt gac cta aga gaa gtg act cgt ttt tta ccc 2158
 Lys Gly Gln Glu Lys Ser Asp Leu Arg Glu Val Thr Arg Phe Leu Pro
 610 615 620 625

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870 875 880	
cca att gct att atc tgg att act ttg aca ttg ttt tat ctt aat tta	2974
Pro Ile Ala Ile Ile Trp Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu	
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Gly His Ile Ser Trp Arg Leu Ser Phe Trp Phe Ile Leu Leu Leu Leu	
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Ser Trp Glu Glu Arg Ile Lys Asp Gly Ile Ile Ile Val Ser Leu Met	
930 935 940 945	
gga gtt cta ttt tat att gca gga cta cta ttc cct atc agg gct cat	3166
Gly Val Leu Phe Tyr Ile Ala Gly Leu Leu Phe Pro Ile Arg Ala His	
950 955 960	
att aca ggt ggt agt att gaa cgc ctg cat tat atc ata gca tgg gag	3214
Ile Thr Gly Gly Ser Ile Glu Arg Leu His Tyr Ile Ile Ala Trp Glu	
965 970 975	
ccg ata gca ttg gct acg ttg att ctt act ctc gtt tat tta tgt ttg	3262
Pro Ile Ala Leu Ala Thr Leu Ile Leu Thr Leu Val Tyr Leu Cys Leu	
980 985 990	
gtt aag att tta caa gga aaa tct tgt cag att ggt gat gtg ttc aat	3310
Val Lys Ile Leu Gln Gly Lys Ser Cys Gln Ile Gly Asp Val Phe Asn	
995 1000 1005	
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Val Asp Arg Tyr Lys Lys Leu Leu Gln Ala Tyr Gly Gly Ser Ser Asp	
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agc ggt tta gcc ttt tta aat gat aaa agg ctc tac tgg tac caa aaa	3406
Ser Gly Leu Ala Phe Leu Asn Asp Lys Arg Leu Tyr Trp Tyr Gln Lys	
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Cys Leu Ile Met Gly Glu Pro Ala Gly Asp Asp Thr Tyr Ile Arg Glu	
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Asp Tyr Phe Gln Gln Ala Pro Ile Ala Leu Val Lys Asn Ala Glu His			
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gaa gtt gtt gct ttt gct aat att atg cca aac tat gaa aag agt att		3934	
Glu Val Val Ala Phe Ala Asn Ile Met Pro Asn Tyr Glu Lys Ser Ile			
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Ile Ser Ile Asp Leu Met Arg His Asp Lys Gln Lys Ile Pro Asn Gly			
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1235	1240	1245	
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Gly Tyr His Tyr Phe Asp Leu Gly Met Ala Pro Leu Ser Gly Val Gly			
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 Lys Lys Phe Thr Pro Leu Trp Ser Glu Arg Tyr Ile Ser Cys Ser Arg
 1300 1305 1310

tcg tcc tgg tta att tgt gct att tgt gcc cta tta atg gaa gat agt 4270
 Ser Ser Trp Leu Ile Cys Ala Ile Cys Ala Leu Leu Met Glu Asp Ser
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 Lys Ile Lys Ile Val Lys
 1330 1335

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 aatatgtgt gcttttaat attgttagc tactgttagt gctgattttt gctttacagc 4438
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 Ala Val Gln Phe Ile Gly Leu Lys Pro Asp Tyr Pro Gly Lys Thr Tyr
 35 40 45
 Phe Ile Ile Leu Leu Thr Ala Trp Thr Leu Met Ala Leu Val Thr Ala
 50 55 60
 Leu Val Gly Trp Asp Asn Arg Tyr Gly Ser Phe Leu Ser Leu Leu Ile
 65 70 75 80
 Leu Leu Phe Gln Leu Gly Ser Ser Ala Gly Thr Tyr Pro Ile Glu Leu
 85 90 95
 Ser Pro Lys Phe Phe Gln Thr Ile Gln Pro Phe Leu Pro Met Thr Tyr
 100 105 110
 Ser Val Ser Gly Leu Arg Glu Thr Ile Ser Leu Thr Gly Asp Val Asn
 115 120 125
 His Gln Trp Arg Met Leu Val Ile Phe Leu Val Ser Ser Met Ile Leu
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 Ile Lys Leu Phe Lys Asn Gln Gly Val Tyr Asn Gly Leu Ile Gly Leu
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 Phe Leu Leu Tyr Gly Leu Tyr Ile Ser Gln Asn Gln Glu Ile Val Ala
 65 70 75 80
 Val Phe Leu Ile Asn Val Leu Leu Val Ala Ile Tyr Gly Ala Leu Thr
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 Lys Leu Asn Thr Ile Ser Phe Arg Gln Ile Ala Val Asp Leu Lys Asp
 65 70 75 80
 Ile Leu Val His Leu Glu Ile Asp Lys Val Ile Leu Val Gly His Ser
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 100 105 110
 Val Arg Gly Leu Leu Leu Asn Ser Gly Asn Leu Thr Ile His Gly Gln
 115 120 125
 Arg Trp Trp Asp Ile Leu Leu Val Arg Ile Ala Tyr Lys Phe Leu His
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 145 150 155 160
 Ser Leu Met Leu Glu Asp Leu Lys Ile Ser Pro Ala Asp Leu Gln His
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 Val Ser Thr Pro Val Met Val Leu Val Gly Asn Lys Asp Ile Ile Lys
 180 185 190
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 195 200 205
 Tyr Ser Leu Val Gly Phe Gly His His Ile Ile Lys Gln Asp Ser His
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 Leu Leu Arg Thr Asp Lys Ser Lys Arg Tyr Ile Leu Gln Thr Ser Trp
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 Cys Ile Asn Thr Phe Asn Asn Leu Ser Gly Phe Gly Gly Leu Ile Asp
 65 70 75 80
 Ile Gly Leu Arg Met Ala Phe Tyr Gly Lys Lys Gly Gln Glu Lys Ser
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 Asp Leu Arg Glu Val Thr Arg Phe Leu Pro Tyr Leu Ile Ser Gly Leu
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 Ser Phe Ile Ser Val Ile Ala Leu Ile Met Ser His Ile Phe His Ala
 115 120 125
 Lys Ala Ser Val Asp Tyr Tyr Leu Val Leu Ile Gly Ala Ser Met
 130 135 140
 Tyr Phe Pro Val Ile Tyr Trp Ile Ser Gly His Lys Gly Ser His Tyr
 145 150 155 160
 Phe Gly Asp Met Pro Ser Ser Thr Arg Ile Lys Leu Gly Val Val Ser
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 Phe Phe Glu Trp Gly Cys Ala Ala Ala Phe Ile Ile Ile Gly Tyr
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 Ile Gly Cys Ala Val Gly Ile Val Ser Leu Ile Pro Gly Gly Leu Gly
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 Ser Phe Glu Leu Val Leu Phe Thr Gly Phe Ala Ala Glu Gly Leu Pro
 225 230 235 240
 Lys Glu Thr Val Val Ala Trp Leu Leu Leu Tyr Arg Leu Ala Tyr Tyr
 245 250 255
 Ile Ile Pro Phe Phe Ala Gly Ile Tyr Phe Phe Ile His Tyr Leu Gly
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 Ser Gln Ile Asn Gln Arg Tyr Glu Asn Val Pro Lys Glu Leu Val Ser
 275 280 285
 Thr Val Leu Gln Thr Met Val Ser His Leu Met Arg Ile Leu Gly Ala
 290 295 300
 Phe Leu Ile Phe Ser Thr Ala Phe Phe Glu Asn Ile Thr Tyr Ile Met
 305 310 315 320
 Trp Leu Gln Lys Leu Gly Leu Asp Pro Leu Gln Glu Gln Met Leu Trp
 325 330 335
 Gln Phe Pro Gly Leu Leu Leu Gly Val Cys Phe Ile Leu Leu Ala Arg
 340 345 350
 Thr Ile Asp Gln Lys Val Lys Asn Ala Phe Pro Ile Ala Ile Ile Trp
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 Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu Gly His Ile Ser Trp Arg
 370 375 380
 Leu Ser Phe Trp Phe Ile Leu Leu Leu Gly Leu Leu Val Ile Lys
 385 390 395 400
 Pro Thr Leu Tyr Lys Lys Gln Phe Ile Tyr Ser Trp Glu Glu Arg Ile
 405 410 415
 Lys Asp Gly Ile Ile Ile Val Ser Leu Met Gly Val Leu Phe Tyr Ile
 420 425 430

Ala Gly Leu Leu Phe Pro Ile Arg Ala His Ile Thr Gly Gly Ser Ile
 435 440 445
 Glu Arg Leu His Tyr Ile Ile Ala Trp Glu Pro Ile Ala Leu Ala Thr
 450 455 460
 Leu Ile Leu Thr Leu Val Tyr Leu Cys Leu Val Lys Ile Leu Gln Gly
 465 470 475 480
 Lys Ser Cys Gln Ile Gly Asp Val Phe Asn Val Asp Arg Tyr Lys Lys
 485 490 495
 Leu Leu Gln Ala Tyr Gly Gly Ser Ser Asp Ser Gly Leu Ala Phe Leu
 500 505 510
 Asn Asp Lys Arg Leu Tyr Trp Tyr Gln Lys Asn Gly Glu Asp Cys Val
 515 520 525
 Ala Phe Gln Phe Val Ile Val Asn Asn Lys Cys Leu Ile Met Gly Glu
 530 535 540
 Pro Ala Gly Asp Asp Thr Tyr Ile Arg Glu Ala Ile Glu Ser Phe Ile
 545 550 555 560
 Asp Asp Ala Asp Lys Leu Asp Tyr Asp Leu Val Phe Tyr Ser Ile Gly
 565 570 575
 Gln Lys Leu Thr Leu Leu His Glu Tyr Gly Phe Asp Phe Met Lys
 580 585 590
 Val Gly Glu Asp Ala Leu Val Asn Leu Glu Thr Phe Thr Leu Lys Gly
 595 600 605
 Asn Lys Tyr Lys Pro Phe Arg Asn Ala Leu Asn Arg Val Glu Lys Asp
 610 615 620
 Gly Phe Tyr Phe Glu Val Val Gln Ser Pro His Ser Gln Glu Leu Leu
 625 630 635 640
 Asn Ser Leu Glu Glu Ile Ser Asn Thr Trp Leu Glu Gly Arg Pro Glu
 645 650 655
 Lys Gly Phe Ser Leu Gly Tyr Phe Asn Lys Asp Tyr Phe Gln Gln Ala
 660 665 670
 Pro Ile Ala Leu Val Lys Asn Ala Glu His Glu Val Val Ala Phe Ala
 675 680 685
 Asn Ile Met Pro Asn Tyr Glu Lys Ser Ile Ile Ser Ile Asp Leu Met
 690 695 700
 Arg His Asp Lys Gln Lys Ile Pro Asn Gly Val Met Asp Phe Leu Phe
 705 710 715 720
 Leu Ser Leu Phe Ser Tyr Tyr Gln Glu Lys Gly Tyr His Tyr Phe Asp
 725 730 735
 Leu Gly Met Ala Pro Leu Ser Gly Val Gly Arg Val Glu Thr Ser Phe
 740 745 750
 Ala Lys Glu Arg Met Ala Tyr Leu Val Tyr His Phe Gly Ser His Phe
 755 760 765
 Tyr Ser Phe Asn Gly Leu His Lys Tyr Lys Lys Phe Thr Pro Leu
 770 775 780
 Trp Ser Glu Arg Tyr Ile Ser Cys Ser Arg Ser Ser Trp Leu Ile Cys
 785 790 795 800
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 805 810 815

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 Asn Ile Thr Tyr Ile Met Trp Leu Gln Lys Leu Gly Leu Asp Pro Leu
 20 25 30
 Gln Glu Gln Met Leu Trp Gln Phe Pro Gly Leu Leu Leu Gly Val Cys
 35 40 45
 Phe Ile Leu Leu Ala Arg Thr Ile Asp Gln Lys Val Lys Asn Ala Phe
 50 55 60
 Pro Ile Ala Ile Ile Trp Ile Thr Leu Thr Leu Phe Tyr Leu Asn Leu
 65 70 75 80
 Gly His Ile Ser Trp Arg Leu Ser Phe Trp Phe Ile Leu Leu Leu Leu
 85 90 95
 Gly Leu Leu Val Ile Lys Pro Thr Leu Tyr Lys Lys Gln Phe Ile Tyr
 100 105 110
 Ser Trp Glu Glu Arg Ile Lys Asp Gly Ile Ile Ile Val Ser Leu Met
 115 120 125
 Gly Val Leu Phe Tyr Ile Ala Gly Leu Leu Phe Pro Ile Arg Ala His
 130 135 140
 Ile Thr Gly Gly Ser Ile Glu Arg Leu His Tyr Ile Ile Ala Trp Glu
 145 150 155 160
 Pro Ile Ala Leu Ala Thr Leu Ile Leu Thr Leu Val Tyr Leu Cys Leu
 165 170 175
 Val Lys Ile Leu Gln Gly Lys Ser Cys Gln Ile Gly Asp Val Phe Asn
 180 185 190
 Val Asp Arg Tyr Lys Lys Leu Leu Gln Ala Tyr Gly Gly Ser Ser Asp
 195 200 205
 Ser Gly Leu Ala Phe Leu Asn Asp Lys Arg Leu Tyr Trp Tyr Gln Lys
 210 215 220
 Asn Gly Glu Asp Cys Val Ala Phe Gln Phe Val Ile Val Asn Asn Lys
 225 230 235 240
 Cys Leu Ile Met Gly Glu Pro Ala Gly Asp Asp Thr Tyr Ile Arg Glu
 245 250 255
 Ala Ile Glu Ser Phe Ile Asp Asp Ala Asp Lys Leu Asp Tyr Asp Leu
 260 265 270
 Val Phe Tyr Ser Ile Gly Gln Lys Leu Thr Leu Leu Leu His Glu Tyr
 275 280 285
 Gly Phe Asp Phe Met Lys Val Gly Glu Asp Ala Leu Val Asn Leu Glu
 290 295 300
 Thr Phe Thr Leu Lys Gly Asn Lys Tyr Lys Pro Phe Arg Asn Ala Leu
 305 310 315 320
 Asn Arg Val Glu Lys Asp Gly Phe Tyr Phe Glu Val Val Gln Ser Pro
 325 330 335
 His Ser Gln Glu Leu Leu Asn Ser Leu Glu Glu Ile Ser Asn Thr Trp
 340 345 350
 Leu Glu Gly Arg Pro Glu Lys Gly Phe Ser Leu Gly Tyr Phe Asn Lys
 355 360 365
 Asp Tyr Phe Gln Gln Ala Pro Ile Ala Leu Val Lys Asn Ala Glu His
 370 375 380
 Glu Val Val Ala Phe Ala Asn Ile Met Pro Asn Tyr Glu Lys Ser Ile
 385 390 395 400
 Ile Ser Ile Asp Leu Met Arg His Asp Lys Gln Lys Ile Pro Asn Gly
 405 410 415
 Val Met Asp Phe Leu Phe Leu Ser Leu Phe Ser Tyr Tyr Gln Glu Lys
 420 425 430
 Gly Tyr His Tyr Phe Asp Leu Gly Met Ala Pro Leu Ser Gly Val Gly
 435 440 445

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Arg Val Glu Thr Ser Phe Ala Lys Glu Arg Met Ala Tyr Leu Val Tyr
 450 455 460
 His Phe Gly Ser His Phe Tyr Ser Phe Asn Gly Leu His Lys Tyr Lys
 465 470 475 480
 Lys Lys Phe Thr Pro Leu Trp Ser Glu Arg Tyr Ile Ser Cys Ser Arg
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 Ser Ser Trp Leu Ile Cys Ala Ile Cys Ala Leu Leu Met Glu Asp Ser
 500 505 510
 Lys Ile Lys Ile Val Lys
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tca tta ttg gag aaa ata tct gtt gag cgt tct ttt att gaa ttt gat 96
 Ser Leu Leu Glu Lys Ile Ser Val Glu Arg Ser Phe Ile Glu Phe Asp
 20 25 30

aaa ctt cta tta gca cct tat tgg cgt aaa gga atg ctg gca cta ata 144
 Lys Leu Leu Leu Ala Pro Tyr Trp Arg Lys Gly Met Leu Ala Leu Ile
 35 40 45

gat agt cat gct ttt aat tat cta cca tgc tta aaa aat agg gaa tta 192
 Asp Ser His Ala Phe Asn Tyr Leu Pro Cys Leu Lys Asn Arg Glu Leu
 50 55 60

caa tta agc gcc ttt ttg tcc cag tta gat aaa gat ttt tta ttt gag 240
 Gln Leu Ser Ala Phe Leu Ser Gln Leu Asp Lys Asp Phe Leu Phe Glu
 65 70 75 80

aca tca gaa caa gct tgg gca tca ctc atc ttg agt atg gaa gtt gaa 288
 Thr Ser Glu Gln Ala Trp Ala Ser Leu Ile Leu Ser Met Glu Val Glu
 85 90 95

cac aca aag act ttt tta aaa aaa tgg aag aca tca act cac ttt caa His Thr Lys Thr Phe Leu Lys Lys Trp Lys Thr Ser Thr His Phe Gln	336
100 105 110	
aaa gat gtt gag cat ata gtg gat gtt tat cgt att cgt gaa caa atg Lys Asp Val Glu His Ile Val Asp Val Tyr Arg Ile Arg Glu Gln Met	384
115 120 125	
gga ttg gct aaa gaa cat ctt tat cgt tat gga aaa act ata ata aaa Gly Leu Ala Lys Glu His Leu Tyr Arg Tyr Gly Lys Thr Ile Ile Lys	432
130 135 140	
caa gcg gaa ggt att cgc aaa gca aga ggc ttg atg gtt gat ttc gaa Gln Ala Glu Gly Ile Arg Lys Ala Arg Gly Leu Met Val Asp Phe Glu	480
145 150 155 160	
aaa ata gaa caa cta gat agt gag tta gca atc cat gat agg cat gag Lys Ile Glu Gln Leu Asp Ser Glu Leu Ala Ile His Asp Arg His Glu	528
165 170 175	
ata gtt gtc aat ggt ggc acc tta atc aag aaa tta gga ata aaa cct Ile Val Val Asn Gly Gly Thr Leu Ile Lys Lys Leu Gly Ile Lys Pro	576
180 185 190	
ggt cca cag atg gga gat att atc tct caa att gaa tta gcc att gtt Gly Pro Gln Met Gly Asp Ile Ile Ser Gln Ile Glu Leu Ala Ile Val	624
195 200 205	
tta gga caa ctg att aat gaa gaa gag gct att tta cat ttt gtt aag Leu Gly Gln Leu Ile Asn Glu Glu Ala Ile Leu His Phe Val Lys	672
210 215 220	
cag tac ttg atg gat tagagaggat tat atg agc gat ttt tta gta gat Gln Tyr Leu Met Asp Met Ser Asp Phe Leu Val Asp	721
225 230 235	
gga ttg act aag tcg gtt ggt gat aag acg gtc ttt agt aat gtt tca Gly Leu Thr Lys Ser Val Gly Asp Lys Thr Val Phe Ser Asn Val Ser	769
240 245 250	
ttt atc atc cat agt tta gac cgt att ggg att att ggt gtc aat gga Phe Ile Ile His Ser Leu Asp Arg Ile Gly Ile Ile Gly Val Asn Gly	817
255 260 265	
act gga aag aca aca cta tta gat gtt att tcg ggt gaa tta ggt ttt Thr Gly Lys Thr Thr Leu Leu Asp Val Ile Ser Gly Glu Leu Gly Phe	865
270 275 280	
gat ggt gat cgt tcc cct ttt tca tca gct aat gat tat aag att gct Asp Gly Asp Arg Ser Pro Phe Ser Ser Ala Asn Asp Tyr Lys Ile Ala	913
285 290 295 300	
tat tta aaa caa gaa cca gac ttt gat gat tct cag aca att ttg gac Tyr Leu Lys Gln Glu Pro Asp Phe Asp Asp Ser Gln Thr Ile Leu Asp	961
305 310 315	

acc gta ctt tct tct gac tta aga gag atg gct tta att aaa gaa tat	320	325	330	1009
Thr Val Leu Ser Ser Asp Leu Arg Glu Met Ala Leu Ile Lys Glu Tyr				
gaa tta ttg ctt aat cac tac gaa gaa agt aag caa tca cgt cta gag	335	340	345	1057
Glu Leu Leu Leu Asn His Tyr Glu Glu Ser Lys Gln Ser Arg Leu Glu				
aaa gta atg gca gaa atg gat tct tta gat gct ttg tct att gag agc	350	355	360	1105
Lys Val Met Ala Glu Met Asp Ser Leu Asp Ala Trp Ser Ile Glu Ser				
gaa gtc aaa aca gta tta tcc aaa tta ggt att act gat ttg cag ttg	365	370	375	1153
Glu Val Lys Thr Val Leu Ser Lys Leu Gly Ile Thr Asp Leu Gln Leu				
tcg gtt ggt gaa tta tca gga gga tta cga aga cgt gtt caa tta gcg	385	390	395	1201
Ser Val Gly Glu Leu Ser Gly Leu Arg Arg Arg Val Gln Leu Ala				
caa gta tta tta aat gat gca gat tta ttg ctc tta gac gaa cct act	400	405	410	1249
Gln Val Leu Leu Asn Asp Ala Asp Leu Leu Leu Asp Glu Pro Thr				
aac cac tta gat att gac act att gca tgg tta acg aat ttt ttg aaa	415	420	425	1297
Asn His Leu Asp Ile Asp Thr Ile Ala Trp Leu Thr Asn Phe Leu Lys				
aat agt aaa aag aca gtg ctt ttt ata act cat gat cgt tat ttt cta	430	435	440	1345
Asn Ser Lys Lys Thr Val Leu Phe Ile Thr His Asp Arg Tyr Phe Leu				
gac aat gtt gca aca cgt att ttt gaa tta gat aag gca cag att aca	445	450	455	1393
Asp Asn Val Ala Thr Arg Ile Phe Glu Leu Asp Lys Ala Gln Ile Thr				
gaa tat caa ggc aat tat cag gat tat gtc cga ctt cgt gca gaa caa	465	470	475	1441
Glu Tyr Gln Gly Asn Tyr Gln Asp Tyr Val Arg Leu Arg Ala Glu Gln				
gac gag cgt gat gct agt tta cat aaa aag aaa cag ctt tat aaa	480	485	490	1489
Asp Glu Arg Asp Ala Ala Ser Leu His Lys Lys Lys Gln Leu Tyr Lys				
cag gaa cta gct tgg atg cgt act cag cca caa gct cgt gca acg aaa	495	500	505	1537
Gln Glu Leu Ala Trp Met Arg Thr Gln Pro Gln Ala Arg Ala Thr Lys				
caa cag gct cgt att aat cgt ttt caa aat cta aaa aac gat tta cac	510	515	520	1585
Gln Gln Ala Arg Ile Asn Arg Phe Gln Asn Leu Lys Asn Asp Leu His				
caa aca agc gat aca agc gat ttg gaa atg aca ttt gaa aca agt cga	525	530	535	1633
Gln Thr Ser Asp Thr Ser Asp Leu Glu Met Thr Phe Glu Thr Ser Arg				

att ggg aaa aag gtt att aat ttt gaa aat gtc tct ttt tct tac cca Ile Gly Lys Lys Val Ile Asn Phe Glu Asn Val Ser Phe Ser Tyr Pro 545 550 555	1681
gat aaa tct atc ttg aaa gac ttt aat ttg tta att caa aat aaa gac Asp Lys Ser Ile Leu Lys Asp Phe Asn Leu Leu Ile Gln Asn Lys Asp 560 565 570	1729
cgt att ggc atc gtt gga gat aat ggt gtt gga aag tca acc tta ctt Arg Ile Gly Ile Val Gly Asp Asn Gly Val Gly Lys Ser Thr Leu Leu 575 580 585	1777
aat tta att gtt caa gat tta cag ccg gat tcg ggt aat gtc tct att Asn Leu Ile Val Gln Asp Leu Gln Pro Asp Ser Gly Asn Val Ser Ile 590 595 600	1825
ggt gaa acg ata cgt gta ggt tac ttt tca caa caa ctt cat aat atg Gly Glu Thr Ile Arg Val Gly Tyr Phe Ser Gln Gln Leu His Asn Met 605 610 615 620	1873
gat ggc tca aaa cgt gtt att aat tat ttg caa gag gtt gca gat gag Asp Gly Ser Lys Arg Val Ile Asn Tyr Leu Gln Glu Val Ala Asp Glu 625 630 635	1921
gtt aaa act agt gtc ggt aca aca agt gtg aca gaa cta ttg gaa caa Val Lys Thr Ser Val Gly Thr Ser Val Thr Glu Leu Leu Glu Gln 640 645 650	1969
ttt ctc ttt cca cgt tcg aca cat gga aca caa att gca aaa tta tca Phe Leu Phe Pro Arg Ser Thr His Gly Thr Gln Ile Ala Lys Leu Ser 655 660 665	2017
ggt ggt gag aaa aaa aga ctt tac ctt tta aaa atc ctg att gaa aag Gly Gly Glu Lys Lys Arg Leu Tyr Leu Leu Lys Ile Leu Ile Glu Lys 670 675 680	2065
cct aat gtg tta cta ctt gat gag ccg aca aat gac tta gat att gct Pro Asn Val Leu Leu Asp Glu Pro Thr Asn Asp Leu Asp Ile Ala 685 690 695 700	2113
aca tta act gtt ctt gaa aat ttt tta caa ggc ttt ggt ggt cct gtg Thr Leu Thr Val Leu Glu Asn Phe Leu Gln Gly Phe Gly Gly Pro Val 705 710 715	2161
att aca gtt agt cac gat cgt tac ttt tta gat aaa gtg gct aat aaa Ile Thr Val Ser His Asp Arg Tyr Phe Leu Asp Lys Val Ala Asn Lys 720 725 730	2209
att att gcg ttt gaa gat aac gat atc cgt gaa ttt ttt ggt aat tat Ile Ile Ala Phe Glu Asp Asn Asp Ile Arg Glu Phe Phe Gly Asn Tyr 735 740 745	2257
act gat tat tta gat gaa aaa gca ttt aat gag caa aat aat gaa gtt Thr Asp Tyr Leu Asp Glu Lys Ala Phe Asn Glu Gln Asn Asn Glu Val 750 755 760	2305

atc agt aaa aaa gag agt acc aag aca agt cgt gaa aag caa agt cgt Ile Ser Lys Lys Glu Ser Thr Lys Thr Ser Arg Glu Lys Gln Ser Arg	765	770	775	780	2353
aaa aga atg tct tac ttt gaa aaa caa gaa tgg gcg aca att gaa gac Lys Arg Met Ser Tyr Phe Glu Lys Gln Glu Trp Ala Thr Ile Glu Asp	785	790	795		2401
gat att atg ata ttg gaa aat act atc act cgt ata gaa aat gat atg Asp Ile Met Ile Leu Glu Asn Thr Ile Thr Arg Ile Glu Asn Asp Met	800	805	810		2449
caa aca tgt ggt agt gat ttt aca agg tta tct gat tta caa aag gaa Gln Thr Cys Gly Ser Asp Phe Thr Arg Leu Ser Asp Leu Gln Lys Glu	815	820	825		2497
tta gat gca aaa aat gaa gca ctt cta gaa aag tat gac cgt tat gag Leu Asp Ala Lys Asn Glu Ala Leu Leu Glu Lys Tyr Asp Arg Tyr Glu	830	835	840		2545
tac ctt agt gag ttagacac atg att atc cgt ccg att att aaa aat gat Tyr Leu Ser Glu LeuAspThrMet Ile Ile Arg Pro Ile Ile Lys Asn Asp	845	850	855	860	2595
gac caa gca gtt gca caa tta att cga caa agt tta cgc gcc tat gat Asp Gln Ala Val Ala Gln Leu Ile Arg Gln Ser Leu Arg Ala Tyr Asp	865	870	875		2643
tta gat aaa cct gat aca gca tat tca gac cct cac tta gat cat ttg Leu Asp Lys Pro Asp Thr Ala Tyr Ser Asp Pro His Leu Asp His Leu	880	885	890		2691
acc tca tac tac gaa aaa ata gag aag tca gga ttc ttt gtc att gag Thr Ser Tyr Tyr Glu Lys Ile Glu Lys Ser Gly Phe Phe Val Ile Glu	895	900	905		2739
gag aga gat gag att att ggc tgt ggc ggc ttt ggt ccg ctg aaa aat Glu Arg Asp Glu Ile Ile Gly Cys Gly Phe Gly Pro Leu Lys Asn	910	915	920	925	2787
cta att gca gag atg cag aag gtg tac att gca gaa cgt ttc cgt ggt Leu Ile Ala Glu Met Gln Lys Val Tyr Ile Ala Glu Arg Phe Arg Gly	930	935	940		2835
aag ggg ctt gct act gat tta gtg aaa atg att gaa gta gaa gct cga Lys Gly Leu Ala Thr Asp Leu Val Lys Met Ile Glu Val Glu Ala Arg	945	950	955		2883
aaa att ggg tat aga caa ctt tat tta gag aca gcc agt act ttg agt Lys Ile Gly Tyr Arg Gln Leu Tyr Leu Glu Thr Ala Ser Thr Leu Ser	960	965	970		2931
agg gca act gcg gtt tat aag cat atg gga tat tgt gcc tta tcg caa Arg Ala Thr Ala Val Tyr Lys His Met Gly Tyr Cys Ala Leu Ser Gln	975	980	985		2979

cca ata gca aat gat caa ggt cat aca gct atg gat att tgg atg att Pro Ile Ala Asn Asp Gln Gly His Thr Ala Met Asp Ile Trp Met Ile 990 995 1000 1005	3027
aaa gat tta taagttgaaa gtggattagt gaacatggat taattatTTT Lys Asp Leu	3076
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aaa agg tac ccc aat tgg tta tgg ctt gat tta cta gga gct atg ctt Lys Arg Tyr Pro Asn Trp Leu Trp Leu Asp Leu Leu Gly Ala Met Leu 1020 1025 1030	3177
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gat aat ggc gtt aca aaa ggt gat cgg act gga gtt tat ctg tgg acg Asp Asn Gly Val Thr Lys Gly Asp Arg Thr Gly Val Tyr Leu Trp Thr 1050 1055 1060	3273
ttc atc atg ttt ata ttt gtt gta cta ggt att att ggg cgt att acg Phe Ile Met Phe Ile Phe Val Val Leu Gly Ile Ile Gly Arg Ile Thr 1065 1070 1075 1080	3321
atg gct tac gca tct agt cgc tta acg aca aca atg att aga gat atg Met Ala Tyr Ala Ser Ser Arg Leu Thr Thr Met Ile Arg Asp Met 1085 1090 1095	3369
cgt aat gat atg tat gct aag ctt caa gaa tac tcc cat cat gaa tat Arg Asn Asp Met Tyr Ala Lys Leu Gln Glu Tyr Ser His His Glu Tyr 1100 1105 1110	3417
gaa cag ata ggt gta tct tca cta gtg aca cgt atg aca agc gat act Glu Gln Ile Gly Val Ser Ser Leu Val Thr Arg Met Thr Ser Asp Thr 1115 1120 1125	3465
ttt gtt ttg atg caa ttt gct gaa atg tct tta cgt tta ggc cta gta Phe Val Leu Met Gln Phe Ala Glu Met Ser Leu Arg Leu Gly Leu Val 1130 1135 1140	3513
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cca tct ttg gct tgg ctt gta gcg gtt gcg atg cct ctt ttg gta gga Pro Ser Leu Ala Trp Leu Val Ala Val Ala Met Pro Leu Leu Val Gly 1165 1170 1175	3609
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aaa ttt caa gtc gct aac caa cgt tac aca gat act tca act ggt ctt Lys Phe Gln Val Ala Asn Gln Arg Tyr Thr Asp Thr Ser Thr Gly Leu 1225 1230 1235 1240	3801
ttt aaa tta aca ggg cta aca gaa cca ctt ttc gtt caa att att att Phe Lys Leu Thr Gly Leu Thr Glu Pro Leu Phe Val Gln Ile Ile Ile 1245 1250 1255	3849
gca atg att gtg gct atc gtt tgg ttt gct ttg gat ccc tta caa aga Ala Met Ile Val Ala Ile Val Trp Phe Ala Leu Asp Pro Leu Gln Arg 1260 1265 1270	3897
ggt gct att aaa ata ggg gat tta gtt gct ttt atc gaa tat agc ttc Gly Ala Ile Lys Ile Gly Asp Leu Val Ala Phe Ile Glu Tyr Ser Phe 1275 1280 1285	3945
cat gct ctc ttt tca ttt ttg cta ttt gcc aat ctt ttt act atg tat His Ala Leu Phe Ser Phe Leu Leu Phe Ala Asn Leu Phe Thr Met Tyr 1290 1295 1300	3993
cct cgt atg gtg gta tca agc cat cgt att aga gag gtg atg gat atg Pro Arg Met Val Val Ser Ser His Arg Ile Arg Glu Val Met Asp Met 1305 1310 1315 1320	4041
cca atc tct atc aat cct aat gcc gaa ggt gtt acg gat acg aaa ctt Pro Ile Ser Ile Asn Pro Asn Ala Glu Gly Val Thr Asp Thr Lys Leu 1325 1330 1335	4089
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aca gag agt ccc gtt ttg cat gat att tct ttt aaa gct aag cct gga Thr Glu Ser Pro Val Leu His Asp Ile Ser Phe Lys Ala Lys Pro Gly 1355 1360 1365	4185
gaa aca att gct ttt att ggt tca aca ggt tca gga aaa tct tct ctt Glu Thr Ile Ala Phe Ile Gly Ser Thr Gly Ser Gly Lys Ser Ser Leu 1370 1375 1380	4233
gtt aat ttg att cca cgt ttt tat gat gtg aca ctt gga aaa atc tta Val Asn Leu Ile Pro Arg Phe Tyr Asp Val Thr Leu Gly Lys Ile Leu 1385 1390 1395 1400	4281
gta gat gga gtt gat gta aga gat tat aac ctt aaa tca ctt cgc caa Val Asp Gly Val Asp Val Arg Asp Tyr Asn Leu Lys Ser Leu Arg Gln 1405 1410 1415	4329

aag att gga ttt atc ccc caa aaa gct ctt tta ttt aca ggg aca ata Lys Ile Gly Phe Ile Pro Gln Lys Ala Leu Leu Phe Thr Gly Thr Ile 1420 1425 1430	4377
gga gag aat tta aaa tat gga aaa gct gat gct act att gat gat ctt Gly Glu Asn Leu Lys Tyr Gly Lys Ala Asp Ala Thr Ile Asp Asp Leu 1435 1440 1445	4425
aga caa gcg gtt gat att tct caa gct aaa gag ttt att gag agt cac Arg Gln Ala Val Asp Ile Ser Gln Ala Lys Glu Phe Ile Glu Ser His 1450 1455 1460	4473
caa gaa gcc ttt gaa acg cat tta gct gaa ggt ggg agc aat ctt tct Gln Glu Ala Phe Glu Thr His Leu Ala Glu Gly Ser Asn Leu Ser 1465 1470 1475 1480	4521
ggg ggt caa aaa caa cgg tta tct att gct agg gct gtt gtt aaa gat Gly Gly Gln Lys Gln Arg Leu Ser Ile Ala Arg Ala Val Val Lys Asp 1485 1490 1495	4569
cca gat tta tat att ttt gat gat tca ttt tct gct ctc gat tat aag Pro Asp Leu Tyr Ile Phe Asp Asp Ser Phe Ser Ala Leu Asp Tyr Lys 1500 1505 1510	4617
aca gac gct act tta aga gcg cgt cta aaa gaa gta acc ggt gat tct Thr Asp Ala Thr Leu Arg Ala Arg Leu Lys Glu Val Thr Gly Asp Ser 1515 1520 1525	4665
aca gtt ttg ata gtt gct caa agg gtg ggt acg att atg gat gct gat Thr Val Leu Ile Val Ala Gln Arg Val Gly Thr Ile Met Asp Ala Asp 1530 1535 1540	4713
cag att att gtc ctt gat gaa ggc gaa att gtc ggt cgt ggt acc cac Gln Ile Ile Val Leu Asp Glu Gly Glu Ile Val Gly Arg Gly Thr His 1545 1550 1555 1560	4761
gct caa tta ata gaa aat aat gct att tat cgt gaa atc gct gag tca Ala Gln Leu Ile Glu Asn Asn Ala Ile Tyr Arg Glu Ile Ala Glu Ser 1565 1570 1575	4809
caa ctg aag aac caa aac tta tca gaa gga gag tgattgt atg aga aaa Gln Leu Lys Asn Gln Asn Leu Ser Glu Gly Glu Met Arg Lys 1580 1585 1590	4858
aaa tct gtt ttt ttg aga tta tgg tct tac cta act cgc tac aaa gct Lys Ser Val Phe Leu Arg Leu Trp Ser Tyr Leu Thr Arg Tyr Lys Ala 1595 1600 1605	4906
act ctt ttc tta gcg att ttt ttg aaa gtt tta tct agt ttt atg agt Thr Leu Phe Leu Ala Ile Phe Leu Lys Val Leu Ser Ser Phe Met Ser 1610 1615 1620	4954
gtt ctg gag cct ttt att tta ggg tta gcg ata aca gag ttg act gct Val Leu Glu Pro Phe Ile Leu Gly Leu Ala Ile Thr Glu Leu Thr Ala 1625 1630 1635	5002

aac ctt gtt gat atg gct aag gga gtt tct ggg gca gaa ttg aac gtt 5050
 Asn Leu Val Asp Met Ala Lys Gly Val Ser Gly Ala Glu Leu Asn Val
 1640 1645 1650

cct tat att gct ggt att ttg att att tat ttt ttc aga ggt gtt ttc 5098
 Pro Tyr Ile Ala Gly Ile Leu Ile Ile Tyr Phe Phe Arg Gly Val Phe
 1655 1660 1665 1670

tat gaa tta ggt tct tat ggc tca aat t 5126
 Tyr Glu Leu Gly Ser Tyr Gly Ser Asn
 1675

<210> 8
 <211> 229
 <212> PRT
 <213> Streptococcus

<400> 8
 Asn Phe Asp Ile Glu Thr Thr Phe Glu Ala Met Lys Lys His Ala
 1 5 10 15
 Ser Leu Leu Glu Lys Ile Ser Val Glu Arg Ser Phe Ile Glu Phe Asp
 20 25 30
 Lys Leu Leu Ala Pro Tyr Trp Arg Lys Gly Met Leu Ala Leu Ile
 35 40 45
 Asp Ser His Ala Phe Asn Tyr Leu Pro Cys Leu Lys Asn Arg Glu Leu
 50 55 60
 Gln Leu Ser Ala Phe Leu Ser Gln Leu Asp Lys Asp Phe Leu Phe Glu
 65 70 75 80
 Thr Ser Glu Gln Ala Trp Ala Ser Leu Ile Leu Ser Met Glu Val Glu
 85 90 95
 His Thr Lys Thr Phe Leu Lys Trp Lys Thr Ser Thr His Phe Gln
 100 105 110
 Lys Asp Val Glu His Ile Val Asp Val Tyr Arg Ile Arg Glu Gln Met
 115 120 125
 Gly Leu Ala Lys Glu His Leu Tyr Arg Tyr Gly Lys Thr Ile Ile Lys
 130 135 140
 Gln Ala Glu Gly Ile Arg Lys Ala Arg Gly Leu Met Val Asp Phe Glu
 145 150 155 160
 Lys Ile Glu Gln Leu Asp Ser Glu Leu Ala Ile His Asp Arg His Glu
 165 170 175
 Ile Val Val Asn Gly Gly Thr Leu Ile Lys Lys Leu Gly Ile Lys Pro
 180 185 190
 Gly Pro Gln Met Gly Asp Ile Ile Ser Gln Ile Glu Leu Ala Ile Val
 195 200 205
 Leu Gly Gln Leu Ile Asn Glu Glu Glu Ala Ile Leu His Phe Val Lys
 210 215 220
 Gln Tyr Leu Met Asp
 225

<210> 9
 <211> 622
 <212> PRT
 <213> Streptococcus

<400> 9

Met Ser Asp Phe Leu Val Asp Gly Leu Thr Lys Ser Val Gly Asp Lys
 1 5 10 15
 Thr Val Phe Ser Asn Val Ser Phe Ile Ile His Ser Leu Asp Arg Ile
 20 25 30
 Gly Ile Ile Gly Val Asn Gly Thr Gly Lys Thr Thr Leu Leu Asp Val
 35 40 45
 Ile Ser Gly Glu Leu Gly Phe Asp Gly Asp Arg Ser Pro Phe Ser Ser
 50 55 60
 Ala Asn Asp Tyr Lys Ile Ala Tyr Leu Lys Gln Glu Pro Asp Phe Asp
 65 70 75 80
 Asp Ser Gln Thr Ile Leu Asp Thr Val Leu Ser Ser Asp Leu Arg Glu
 85 90 95
 Met Ala Leu Ile Lys Glu Tyr Glu Leu Leu Asn His Tyr Glu Glu
 100 105 110
 Ser Lys Gln Ser Arg Leu Glu Lys Val Met Ala Glu Met Asp Ser Leu
 115 120 125
 Asp Ala Trp Ser Ile Glu Ser Glu Val Lys Thr Val Leu Ser Lys Leu
 130 135 140
 Gly Ile Thr Asp Leu Gln Leu Ser Val Gly Glu Leu Ser Gly Gly Leu
 145 150 155 160
 Arg Arg Arg Val Gln Leu Ala Gln Val Leu Leu Asn Asp Ala Asp Leu
 165 170 175
 Leu Leu Leu Asp Glu Pro Thr Asn His Leu Asp Ile Asp Thr Ile Ala
 180 185 190
 Trp Leu Thr Asn Phe Leu Lys Asn Ser Lys Lys Thr Val Leu Phe Ile
 195 200 205
 Thr His Asp Arg Tyr Phe Leu Asp Asn Val Ala Thr Arg Ile Phe Glu
 210 215 220
 Leu Asp Lys Ala Gln Ile Thr Glu Tyr Gln Gly Asn Tyr Gln Asp Tyr
 225 230 235 240
 Val Arg Leu Arg Ala Glu Gln Asp Glu Arg Asp Ala Ala Ser Leu His
 245 250 255
 Lys Lys Lys Gln Leu Tyr Lys Gln Glu Leu Ala Trp Met Arg Thr Gln
 260 265 270
 Pro Gln Ala Arg Ala Thr Lys Gln Gln Ala Arg Ile Asn Arg Phe Gln
 275 280 285
 Asn Leu Lys Asn Asp Leu His Gln Thr Ser Asp Thr Ser Asp Leu Glu
 290 295 300
 Met Thr Phe Glu Thr Ser Arg Ile Gly Lys Lys Val Ile Asn Phe Glu
 305 310 315 320
 Asn Val Ser Phe Ser Tyr Pro Asp Lys Ser Ile Leu Lys Asp Phe Asn
 325 330 335
 Leu Leu Ile Gln Asn Lys Asp Arg Ile Gly Ile Val Gly Asp Asn Gly
 340 345 350
 Val Gly Lys Ser Thr Leu Leu Asn Leu Ile Val Gln Asp Leu Gln Pro
 355 360 365
 Asp Ser Gly Asn Val Ser Ile Gly Glu Thr Ile Arg Val Gly Tyr Phe
 370 375 380
 Ser Gln Gln Leu His Asn Met Asp Gly Ser Lys Arg Val Ile Asn Tyr
 385 390 395 400
 Leu Gln Glu Val Ala Asp Glu Val Lys Thr Ser Val Gly Thr Thr Ser
 405 410 415
 Val Thr Glu Leu Leu Glu Gln Phe Leu Phe Pro Arg Ser Thr His Gly
 420 425 430

Thr Gln Ile Ala Lys Leu Ser Gly Gly Glu Lys Lys Arg Leu Tyr Leu
 435 440 445
 Leu Lys Ile Leu Ile Glu Lys Pro Asn Val Leu Leu Asp Glu Pro
 450 455 460
 Thr Asn Asp Leu Asp Ile Ala Thr Leu Thr Val Leu Glu Asn Phe Leu
 465 470 475 480
 Gln Gly Phe Gly Gly Pro Val Ile Thr Val Ser His Asp Arg Tyr Phe
 485 490 495
 Leu Asp Lys Val Ala Asn Lys Ile Ile Ala Phe Glu Asp Asn Asp Ile
 500 505 510
 Arg Glu Phe Phe Gly Asn Tyr Thr Asp Tyr Leu Asp Glu Lys Ala Phe
 515 520 525
 Asn Glu Gln Asn Asn Glu Val Ile Ser Lys Lys Glu Ser Thr Lys Thr
 530 535 540
 Ser Arg Glu Lys Gln Ser Arg Lys Arg Met Ser Tyr Phe Glu Lys Gln
 545 550 555 560
 Glu Trp Ala Thr Ile Glu Asp Asp Ile Met Ile Leu Glu Asn Thr Ile
 565 570 575
 Thr Arg Ile Glu Asn Asp Met Gln Thr Cys Gly Ser Asp Phe Thr Arg
 580 585 590
 Leu Ser Asp Leu Gln Lys Glu Leu Asp Ala Lys Asn Glu Ala Leu Leu
 595 600 605
 Glu Lys Tyr Asp Arg Tyr Glu Tyr Leu Ser Glu Leu Asp Thr
 610 615 620

<210> 10
 <211> 157
 <212> PRT
 <213> Streptococcus

<400> 10
 Met Ile Ile Arg Pro Ile Ile Lys Asn Asp Asp Gln Ala Val Ala Gln
 1 5 10 15
 Leu Ile Arg Gln Ser Leu Arg Ala Tyr Asp Leu Asp Lys Pro Asp Thr
 20 25 30
 Ala Tyr Ser Asp Pro His Leu Asp His Leu Thr Ser Tyr Tyr Glu Lys
 35 40 45
 Ile Glu Lys Ser Gly Phe Phe Val Ile Glu Glu Arg Asp Glu Ile Ile
 50 55 60
 Gly Cys Gly Gly Phe Gly Pro Leu Lys Asn Leu Ile Ala Glu Met Gln
 65 70 75 80
 Lys Val Tyr Ile Ala Glu Arg Phe Arg Gly Lys Gly Leu Ala Thr Asp
 85 90 95
 Leu Val Lys Met Ile Glu Val Glu Ala Arg Lys Ile Gly Tyr Arg Gln
 100 105 110
 Leu Tyr Leu Glu Thr Ala Ser Thr Leu Ser Arg Ala Thr Ala Val Tyr
 115 120 125
 Lys His Met Gly Tyr Cys Ala Leu Ser Gln Pro Ile Ala Asn Asp Gln
 130 135 140
 Gly His Thr Ala Met Asp Ile Trp Met Ile Lys Asp Leu
 145 150 155

<210> 11
 <211> 579
 <212> PRT
 <213> Streptococcus

<400> 11
 Met Ala Tyr Ile Trp Ser Tyr Leu Lys Arg Tyr Pro Asn Trp Leu Trp
 1 5 10 15
 Leu Asp Leu Leu Gly Ala Met Leu Phe Val Thr Val Ile Leu Gly Met
 20 25 30
 Pro Thr Ala Leu Ala Gly Met Ile Asp Asn Gly Val Thr Lys Gly Asp
 35 40 45
 Arg Thr Gly Val Tyr Leu Trp Thr Phe Ile Met Phe Ile Phe Val Val
 50 55 60
 Leu Gly Ile Ile Gly Arg Ile Thr Met Ala Tyr Ala Ser Ser Arg Leu
 65 70 75 80
 Thr Thr Thr Met Ile Arg Asp Met Arg Asn Asp Met Tyr Ala Lys Leu
 85 90 95
 Gln Glu Tyr Ser His His Glu Tyr Glu Gln Ile Gly Val Ser Ser Leu
 100 105 110
 Val Thr Arg Met Thr Ser Asp Thr Phe Val Leu Met Gln Phe Ala Glu
 115 120 125
 Met Ser Leu Arg Leu Gly Leu Val Thr Pro Met Val Met Ile Phe Ser
 130 135 140
 Val Val Met Ile Leu Ile Thr Ser Pro Ser Leu Ala Trp Leu Val Ala
 145 150 155 160
 Val Ala Met Pro Leu Leu Val Gly Val Val Leu Tyr Val Ala Ile Lys
 165 170 175
 Thr Lys Pro Leu Ser Glu Arg Gln Gln Thr Met Leu Asp Lys Ile Asn
 180 185 190
 Gln Tyr Val Arg Glu Asn Leu Thr Gly Leu Arg Val Val Arg Ala Phe
 195 200 205
 Ala Arg Glu Asn Phe Gln Ser Gln Lys Phe Gln Val Ala Asn Gln Arg
 210 215 220
 Tyr Thr Asp Thr Ser Thr Gly Leu Phe Lys Leu Thr Gly Leu Thr Glu
 225 230 235 240
 Pro Leu Phe Val Gln Ile Ile Ile Ala Met Ile Val Ala Ile Val Trp
 245 250 255
 Phe Ala Leu Asp Pro Leu Gln Arg Gly Ala Ile Lys Ile Gly Asp Leu
 260 265 270
 Val Ala Phe Ile Glu Tyr Ser Phe His Ala Leu Phe Ser Phe Leu Leu
 275 280 285
 Phe Ala Asn Leu Phe Thr Met Tyr Pro Arg Met Val Val Ser Ser His
 290 295 300
 Arg Ile Arg Glu Val Met Asp Met Pro Ile Ser Ile Asn Pro Asn Ala
 305 310 315 320
 Glu Gly Val Thr Asp Thr Lys Leu Lys Gly His Leu Glu Phe Asp Asn
 325 330 335
 Val Thr Phe Ala Tyr Pro Gly Glu Thr Glu Ser Pro Val Leu His Asp
 340 345 350
 Ile Ser Phe Lys Ala Lys Pro Gly Glu Thr Ile Ala Phe Ile Gly Ser
 355 360 365
 Thr Gly Ser Gly Lys Ser Ser Leu Val Asn Leu Ile Pro Arg Phe Tyr
 370 375 380
 Asp Val Thr Leu Gly Lys Ile Leu Val Asp Gly Val Asp Val Arg Asp
 385 390 395 400
 Tyr Asn Leu Lys Ser Leu Arg Gln Lys Ile Gly Phe Ile Pro Gln Lys
 405 410 415
 Ala Leu Leu Phe Thr Gly Thr Ile Gly Glu Asn Leu Lys Tyr Gly Lys
 420 425 430

Ala Asp Ala Thr Ile Asp Asp Leu Arg Gln Ala Val Asp Ile Ser Gln
 435 440 445
 Ala Lys Glu Phe Ile Glu Ser His Gln Glu Ala Phe Glu Thr His Leu
 450 455 460
 Ala Glu Gly Ser Asn Leu Ser Gly Gly Gln Lys Gln Arg Leu Ser
 465 470 475 480
 Ile Ala Arg Ala Val Val Lys Asp Pro Asp Leu Tyr Ile Phe Asp Asp
 485 490 495
 Ser Phe Ser Ala Leu Asp Tyr Lys Thr Asp Ala Thr Leu Arg Ala Arg
 500 505 510
 Leu Lys Glu Val Thr Gly Asp Ser Thr Val Leu Ile Val Ala Gln Arg
 515 520 525
 Val Gly Thr Ile Met Asp Ala Asp Gln Ile Ile Val Leu Asp Glu Gly
 530 535 540
 Glu Ile Val Gly Arg Gly Thr His Ala Gln Leu Ile Glu Asn Asn Ala
 545 550 555 560
 Ile Tyr Arg Glu Ile Ala Glu Ser Gln Leu Lys Asn Gln Asn Leu Ser
 565 570 575
 Glu Gly Glu

<210> 12
 <211> 92
 <212> PRT
 <213> Streptococcus

<400> 12
 Met Arg Lys Lys Ser Val Phe Leu Arg Leu Trp Ser Tyr Leu Thr Arg
 1 5 10 15
 Tyr Lys Ala Thr Leu Phe Leu Ala Ile Phe Leu Lys Val Leu Ser Ser
 20 25 30
 Phe Met Ser Val Leu Glu Pro Phe Ile Leu Gly Leu Ala Ile Thr Glu
 35 40 45
 Leu Thr Ala Asn Leu Val Asp Met Ala Lys Gly Val Ser Gly Ala Glu
 50 55 60
 Leu Asn Val Pro Tyr Ile Ala Gly Ile Leu Ile Ile Tyr Phe Phe Arg
 65 70 75 80
 Gly Val Phe Tyr Glu Leu Gly Ser Tyr Gly Ser Asn
 85 90

<210> 13
 <211> 5215
 <212> DNA
 <213> Streptococcus

<220>
 <221> CDS
 <222> (3)...(122)

<221> CDS
 <222> (133)...(2511)

<221> CDS
 <222> (367)...(2511)

<221> CDS

<222> (2946) ... (2716)
 <223> of complementary strand

<221> CDS
 <222> (3252) ... (2995)
 <223> of complementary strand

<221> CDS
 <222> (3676) ... (3299)
 <223> of complementary strand

<221> CDS
 <222> (4124) ... (3837)
 <223> of complementary strand

<221> CDS
 <222> (5214) ... (4351)
 <223> of complementary strand

<400> 13

aa	ttt	gga	agt	gct	cta	tca	aca	gtt	gaa	gta	aag	gag	att	att	att	agt	47
		Phe	Gly	Ser	Ala	Leu	Ser	Thr	Val	Glu	Val	Lys	Glu	Ile	Ile	Ser	
1		5				10						15					

gaa gaa aac ata tgg tta tat cgg ctc agt tgc tgc cat ttt act agc

Glu	Glu	Asn	Ile	Trp	Leu	Tyr	Arg	Leu	Ser	Cys	Cys	His	Phe	Thr	Ser	95
20					25							30				

tac tca tat tgg aag tta cca act tgg taagcatcat atg ggt cta gca

Tyr	Ser	Tyr	Trp	Lys	Leu	Pro	Thr	Trp					Met	Gly	Leu	Ala	144
35				40													

aca aag gac aat cag att gcc tat att gat gac agc aaa ggt aag gca

Thr	Lys	Asp	Asn	Gln	Ile	Ala	Tyr	Ile	Asp	Asp	Ser	Lys	Gly	Lys	Ala	192
45				50				55				60				

aaa gcc cct aaa aca aac aaa acg atg gat caa atc agt gct gaa gaa

Lys	Ala	Pro	Lys	Thr	Asn	Lys	Thr	Met	Asp	Gln	Ile	Ser	Ala	Glu	Glu	240
65				70				75								

ggc atc tct gct gaa cag atc gta gtc aaa att act gac caa ggc tat

Gly	Ile	Ser	Ala	Glu	Gln	Ile	Val	Val	Lys	Ile	Thr	Asp	Gln	Gly	Tyr	288
80				85				90								

gtg acc tca cac ggt gac cat tat cat ttt tac aat ggg aaa gtt cct

Val	Thr	Ser	His	Gly	Asp	His	Tyr	His	Phe	Tyr	Asn	Gly	Lys	Val	Pro	336
95					100			105								

tat gat gcg att att agt gaa gag ttg ttg atg acg gat cct aat tac

Tyr	Asp	Ala	Ile	Ile	Ser	Glu	Glu	Leu	Leu	Met	Thr	Asp	Pro	Asn	Tyr	384
110				115				120								

cgt ttt aaa caa tca gac gtt atc aat gaa atc tta gac ggt tac gtt

Arg	Phe	Lys	Gln	Ser	Asp	Val	Ile	Asn	Glu	Ile	Leu	Asp	Gly	Tyr	Val	432
125					130			135				140				

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att aaa gtc aat ggc aac tat tat gtt tac ctc aag cca ggt agt aag Ile Lys Val Asn Gly Asn Tyr Tyr Val Tyr Leu Lys Pro Gly Ser Lys 145 150 155	480
cgc aaa aac att cga acc aaa caa caa att gct gag caa gta gcc aaa Arg Lys Asn Ile Arg Thr Lys Gln Gln Ile Ala Glu Gln Val Ala Lys 160 165 170	528
gga act aaa gaa gct aaa gaa aaa ggt tta gct caa gtg gcc cat ctc Gly Thr Lys Glu Ala Lys Glu Lys Gly Leu Ala Gln Val Ala His Leu 175 180 185	576
agt aaa gaa gaa gtt gcg gca gtc aat gaa gca aaa aga caa gga cgc Ser Lys Glu Glu Val Ala Ala Val Asn Glu Ala Lys Arg Gln Gly Arg 190 195 200	624
tat act aca gac gat ggc tat att ttt agt ccg aca gat atc att gat Tyr Thr Thr Asp Asp Gly Tyr Ile Phe Ser Pro Thr Asp Ile Ile Asp 205 210 215 220	672
gat tta gga gat gct tat tta gta cct cat ggt aat cac tat cat tat Asp Leu Gly Asp Ala Tyr Leu Val Pro His Gly Asn His Tyr His Tyr 225 230 235	720
att cct aaa aag gat ttg tct cca agt gag cta gct gct gca caa gcc Ile Pro Lys Lys Asp Leu Ser Pro Ser Glu Leu Ala Ala Ala Gln Ala 240 245 250	768
tac tgg agt caa aaa caa ggt cga ggt gct aga ccg tct gat tac cgc Tyr Trp Ser Gln Lys Gln Gly Arg Gly Ala Arg Pro Ser Asp Tyr Arg 255 260 265	816
ccg aca cca gcc cca ggt cgt agg aaa gcc cca att cct gat gtg acg Pro Thr Pro Ala Pro Gly Arg Arg Lys Ala Pro Ile Pro Asp Val Thr 270 275 280	864
cct aac cct gga caa ggt cat cag cca gat aac ggt ggc tat cat cca Pro Asn Pro Gly Gln Gly His Gln Pro Asp Asn Gly Gly Tyr His Pro 285 290 295 300	912
gcg cct cct agg cca aat gat gcg tca caa aac aaa cac caa aga gat Ala Pro Pro Arg Pro Asn Asp Ala Ser Gln Asn Lys His Gln Arg Asp 305 310 315	960
gag ttt aaa gga aaa acc ttt aag gaa ctt tta gat caa cta cac cgt Glu Phe Lys Gly Lys Thr Phe Lys Glu Leu Leu Asp Gln Leu His Arg 320 325 330	1008
ctt gat ttg aaa tac cgt cat gtg gaa gaa gat ggg ttg att ttt gaa Leu Asp Leu Lys Tyr Arg His Val Glu Glu Asp Gly Leu Ile Phe Glu 335 340 345	1056
ccg act caa gtg atc aaa tca aac gct ttt ggg tat gtg gtg cct cat Pro Thr Gln Val Ile Lys Ser Asn Ala Phe Gly Tyr Val Val Pro His 350 355 360	1104

gga gat cat tat cat att atc cca aga agt cag tta tca cct ctt gaa	1152
Gly Asp His Tyr His Ile Ile Pro Arg Ser Gln Leu Ser Pro Leu Glu	
365 370 375 380	
atg gaa tta gca gat cga tac tta gct ggc caa act gag gac aat gac	1200
Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly Gln Thr Glu Asp Asn Asp	
385 390 395	
tca ggt tca gag cac tca aaa cca tca gat aaa gaa gtg aca cat acc	1248
Ser Gly Ser Glu His Ser Lys Pro Ser Asp Lys Glu Val Thr His Thr	
400 405 410	
ttt ctt ggt cat cgc atc aaa gct tac gga aaa ggc tta gat ggt aaa	1296
Phe Leu Gly His Arg Ile Lys Ala Tyr Gly Lys Gly Leu Asp Gly Lys	
415 420 425	
cca tat gat acg agt gat gct tat gtt ttt agt aaa gaa tcc att cat	1344
Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe Ser Lys Glu Ser Ile His	
430 435 440	
tca gtg gat aaa tca gga gtt aca gct aaa cac gga gat cat ttc cac	1392
Ser Val Asp Lys Ser Gly Val Thr Ala Lys His Gly Asp His Phe His	
445 450 455 460	
tat ata gga ttt gga gaa ctt gaa caa tat gag ttg gat gag gtc gct	1440
Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr Glu Leu Asp Glu Val Ala	
465 470 475	
aac tgg gtg aaa gca aaa ggt caa gct gat gag ctt gct gct gct ttg	1488
Asn Trp Val Lys Ala Lys Gly Gln Ala Asp Glu Leu Ala Ala Leu	
480 485 490	
gat cag gaa caa ggc aaa gaa aaa cca ctc ttt gac act aaa aaa gtg	1536
Asp Gln Glu Gln Gly Lys Glu Lys Pro Leu Phe Asp Thr Lys Lys Val	
495 500 505	
agt cgc aaa gta aca aaa gat ggt aaa gtg ggc tat atg atg cca aaa	1584
Ser Arg Lys Val Thr Lys Asp Gly Lys Val Gly Tyr Met Met Pro Lys	
510 515 520	
gat ggt aag gac tat ttc tat gct cgt gat caa ctt gat ttg act cag	1632
Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp Gln Leu Asp Leu Thr Gln	
525 530 535 540	
att gcc ttt gcc gaa caa gaa cta atg ctt aaa gat aag aag cat tac	1680
Ile Ala Phe Ala Glu Gln Glu Leu Met Leu Lys Asp Lys Lys His Tyr	
545 550 555	
cgt tat gac att gtt gac aca ggt att gag cca cga ctt gct gta gat	1728
Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu Pro Arg Leu Ala Val Asp	
560 565 570	
gtg tca agt ctg ccg atg cat gct ggt aat gct act tac gat act gga	1776
Val Ser Ser Leu Pro Met His Ala Gly Asn Ala Thr Tyr Asp Thr Gly	
575 580 585	

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agt tcg ttt gtt atc cca cat att gat cat atc cat gtc gtt ccg tat		1824
Ser Ser Phe Val Ile Pro His Ile Asp His Ile His Val Val Pro Tyr		
590	595	600
tca tgg ttg acg cgc gat cag att gca aca gtc aag tat gtg atg caa		1872
Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr Val Lys Tyr Val Met Gln		
605	610	615
620		
cac ccc gaa gtt cgt ccg gat gta tgg tct aag cca ggg cat gaa gag		1920
His Pro Glu Val Arg Pro Asp Val Trp Ser Lys Pro Gly His Glu Glu		
625	630	635
tca ggt tcg gtc att cca aat gtt acg cct ctt gat aaa cgt gct ggt		1968
Ser Gly Ser Val Ile Pro Asn Val Thr Pro Leu Asp Lys Arg Ala Gly		
640	645	650
atg cca aac tgg caa att atc cat tct gct gaa gaa gtt caa aaa gcc		2016
Met Pro Asn Trp Gln Ile Ile His Ser Ala Glu Glu Val Gln Lys Ala		
655	660	665
cta gca gaa ggt cgt ttt gca aca cca gac ggc tat att ttc gat cca		2064
Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp Gly Tyr Ile Phe Asp Pro		
670	675	680
cga gat gtt ttg gcc aaa gaa act ttt gta tgg aaa gat ggc tcc ttt		2112
Arg Asp Val Leu Ala Lys Glu Thr Phe Val Trp Lys Asp Gly Ser Phe		
685	690	695
700		
agc atc cca aga gca gat ggc agt tca ttg aga acc att aat aaa tct		2160
Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu Arg Thr Ile Asn Lys Ser		
705	710	715
gat cta tcc caa gct gag tgg caa caa gct caa gag tta ttg gca aag		2208
Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala Gln Glu Leu Leu Ala Lys		
720	725	730
aaa aat act ggt gat gct act gat acg gat aaa ccc aaa gaa aag caa		2256
Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp Lys Pro Lys Glu Lys Gln		
735	740	745
cag gca gat aag agc aat gaa aac caa cag cca agt gaa gcc agt aaa		2304
Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln Pro Ser Glu Ala Ser Lys		
750	755	760
gaa gaa aaa gaa tca gat gac ttt ata gac agt tta cca gac tat ggt		2352
Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp Ser Leu Pro Asp Tyr Gly		
765	770	775
780		
ctg gat aga gca acc cta gaa gat cat atc aat caa tta gca caa aaa		2400
Leu Asp Arg Ala Thr Leu Glu Asp His Ile Asn Gln Leu Ala Gln Lys		
785	790	795
gct aat atc gat cct aag tat ctc att ttc caa cca gaa ggt gtc caa		2448
Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe Gln Pro Glu Gly Val Gln		
800	805	810

<213> Streptococcus

<400> 14

Phe	Gly	Ser	Ala	Leu	Ser	Thr	Val	Glu	Val	Lys	Glu	Ile	Ile	Ser	Glu
1				5				10				15			
Glu	Asn	Ile	Trp	Leu	Tyr	Arg	Leu	Ser	Cys	Cys	His	Phe	Thr	Ser	Tyr
				20				25				30			
Ser	Tyr	Trp	Lys	Leu	Pro	Thr	Trp								
				35				40							

<210> 15

<211> 793

<212> PRT

<213> Streptococcus

<400> 15

Met	Gly	Leu	Ala	Thr	Lys	Asp	Asn	Gln	Ile	Ala	Tyr	Ile	Asp	Asp	Ser
1				5				10			15				
Lys	Gly	Lys	Ala	Lys	Ala	Pro	Lys	Thr	Asn	Lys	Thr	Met	Asp	Gln	Ile
								20		25		30			
Ser	Ala	Glu	Glu	Gly	Ile	Ser	Ala	Glu	Gln	Ile	Val	Val	Lys	Ile	Thr
					35			40		45					
Asp	Gln	Gly	Tyr	Val	Thr	Ser	His	Gly	Asp	His	Tyr	His	Phe	Tyr	Asn
				50			55		60						
Gly	Lys	Val	Pro	Tyr	Asp	Ala	Ile	Ile	Ser	Glu	Glu	Leu	Leu	Met	Thr
				65			70		75			80			
Asp	Pro	Asn	Tyr	Arg	Phe	Lys	Gln	Ser	Asp	Val	Ile	Asn	Glu	Ile	Leu
				85			90		95						
Asp	Gly	Tyr	Val	Ile	Lys	Val	Asn	Gly	Asn	Tyr	Tyr	Val	Tyr	Leu	Lys
				100			105		110						
Pro	Gly	Ser	Lys	Arg	Lys	Asn	Ile	Arg	Thr	Lys	Gln	Gln	Ile	Ala	Glu
				115			120		125						
Gln	Val	Ala	Lys	Gly	Thr	Lys	Glu	Ala	Lys	Glu	Lys	Gly	Leu	Ala	Gln
				130			135		140						
Val	Ala	His	Leu	Ser	Lys	Glu	Glu	Val	Ala	Ala	Val	Asn	Glu	Ala	Lys
				145			150		155			160			
Arg	Gln	Gly	Arg	Tyr	Thr	Thr	Asp	Asp	Gly	Tyr	Ile	Phe	Ser	Pro	Thr
				165			170		175						
Asp	Ile	Ile	Asp	Asp	Leu	Gly	Asp	Ala	Tyr	Leu	Val	Pro	His	Gly	Asn
				180			185		190						
His	Tyr	His	Tyr	Ile	Pro	Lys	Lys	Asp	Leu	Ser	Pro	Ser	Glu	Leu	Ala
				195			200		205						
Ala	Ala	Gln	Ala	Tyr	Trp	Ser	Gln	Lys	Gln	Gly	Arg	Gly	Ala	Arg	Pro
				210			215		220						
Ser	Asp	Tyr	Arg	Pro	Thr	Pro	Ala	Pro	Gly	Arg	Arg	Lys	Ala	Pro	Ile
				225			230		235			240			
Pro	Asp	Val	Thr	Pro	Asn	Pro	Gly	Gln	Gly	His	Gln	Pro	Asp	Asn	Gly
				245			250		255						
Gly	Tyr	His	Pro	Ala	Pro	Pro	Arg	Pro	Asn	Asp	Ala	Ser	Gln	Asn	Lys
				260			265		270						
His	Gln	Arg	Asp	Glu	Phe	Lys	Gly	Lys	Thr	Phe	Lys	Glu	Leu	Leu	Asp
				275			280		285						
Gln	Leu	His	Arg	Leu	Asp	Leu	Lys	Tyr	Arg	His	Val	Glu	Glu	Asp	Gly
				290			295		300						
Leu	Ile	Phe	Glu	Pro	Thr	Gln	Val	Ile	Lys	Ser	Asn	Ala	Phe	Gly	Tyr
				305			310		315			320			

Val Val Pro His Gly Asp His Tyr His Ile Ile Pro Arg Ser Gln Leu
 325 330 335
 Ser Pro Leu Glu Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly Gln Thr
 340 345 350
 Glu Asp Asn Asp Ser Gly Ser Glu His Ser Lys Pro Ser Asp Lys Glu
 355 360 365
 Val Thr His Thr Phe Leu Gly His Arg Ile Lys Ala Tyr Gly Lys Gly
 370 375 380
 Leu Asp Gly Lys Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe Ser Lys
 385 390 395 400
 Glu Ser Ile His Ser Val Asp Lys Ser Gly Val Thr Ala Lys His Gly
 405 410 415
 Asp His Phe His Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr Glu Leu
 420 425 430
 Asp Glu Val Ala Asn Trp Val Lys Ala Lys Gly Gln Ala Asp Glu Leu
 435 440 445
 Ala Ala Ala Leu Asp Gln Glu Gln Gly Lys Glu Lys Pro Leu Phe Asp
 450 455 460
 Thr Lys Lys Val Ser Arg Lys Val Thr Lys Asp Gly Lys Val Gly Tyr
 465 470 475 480
 Met Met Pro Lys Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp Gln Leu
 485 490 495
 Asp Leu Thr Gln Ile Ala Phe Ala Glu Gln Glu Leu Met Leu Lys Asp
 500 505 510
 Lys Lys His Tyr Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu Pro Arg
 515 520 525
 Leu Ala Val Asp Val Ser Ser Leu Pro Met His Ala Gly Asn Ala Thr
 530 535 540
 Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His Ile His
 545 550 555 560
 Val Val Pro Tyr Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr Val Lys
 565 570 575
 Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser Lys Pro
 580 585 590
 Gly His Glu Glu Ser Gly Ser Val Ile Pro Asn Val Thr Pro Leu Asp
 595 600 605
 Lys Arg Ala Gly Met Pro Asn Trp Gln Ile Ile His Ser Ala Glu Glu
 610 615 620
 Val Gln Lys Ala Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp Gly Tyr
 625 630 635 640
 Ile Phe Asp Pro Arg Asp Val Leu Ala Lys Glu Thr Phe Val Trp Lys
 645 650 655
 Asp Gly Ser Phe Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu Arg Thr
 660 665 670
 Ile Asn Lys Ser Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala Gln Glu
 675 680 685
 Leu Leu Ala Lys Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp Lys Pro
 690 695 700
 Lys Glu Lys Gln Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln Pro Ser
 705 710 715 720
 Glu Ala Ser Lys Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp Ser Leu
 725 730 735
 Pro Asp Tyr Gly Leu Asp Arg Ala Thr Leu Glu Asp His Ile Asn Gln
 740 745 750
 Leu Ala Gln Lys Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe Gln Pro
 755 760 765

Glu Gly Val Gln Phe Tyr Asn Lys Asn Gly Glu Leu Val Thr Tyr Asp
 770 775 780
 Ile Lys Thr Leu Gln Gln Ile Asn Pro
 785 790

<210> 16
 <211> 715
 <212> PRT
 <213> Streptococcus

<400> 16
 Met Thr Asp Pro Asn Tyr Arg Phe Lys Gln Ser Asp Val Ile Asn Glu
 1 5 10 15
 Ile Leu Asp Gly Tyr Val Ile Lys Val Asn Gly Asn Tyr Tyr Val Tyr
 20 25 30
 Leu Lys Pro Gly Ser Lys Arg Lys Asn Ile Arg Thr Lys Gln Gln Ile
 35 40 45
 Ala Glu Gln Val Ala Lys Gly Thr Lys Glu Ala Lys Glu Lys Gly Leu
 50 55 60
 Ala Gln Val Ala His Leu Ser Lys Glu Glu Val Ala Ala Val Asn Glu
 65 70 75 80
 Ala Lys Arg Gln Gly Arg Tyr Thr Thr Asp Asp Gly Tyr Ile Phe Ser
 85 90 95
 Pro Thr Asp Ile Ile Asp Asp Leu Gly Asp Ala Tyr Leu Val Pro His
 100 105 110
 Gly Asn His Tyr His Tyr Ile Pro Lys Lys Asp Leu Ser Pro Ser Glu
 115 120 125
 Leu Ala Ala Ala Gln Ala Tyr Trp Ser Gln Lys Gln Gly Arg Gly Ala
 130 135 140
 Arg Pro Ser Asp Tyr Arg Pro Thr Pro Ala Pro Gly Arg Arg Lys Ala
 145 150 155 160
 Pro Ile Pro Asp Val Thr Pro Asn Pro Gly Gln Gly His Gln Pro Asp
 165 170 175
 Asn Gly Gly Tyr His Pro Ala Pro Pro Arg Pro Asn Asp Ala Ser Gln
 180 185 190
 Asn Lys His Gln Arg Asp Glu Phe Lys Gly Lys Thr Phe Lys Glu Leu
 195 200 205
 Leu Asp Gln Leu His Arg Leu Asp Leu Lys Tyr Arg His Val Glu Glu
 210 215 220
 Asp Gly Leu Ile Phe Glu Pro Thr Gln Val Ile Lys Ser Asn Ala Phe
 225 230 235 240
 Gly Tyr Val Val Pro His Gly Asp His Tyr His Ile Ile Pro Arg Ser
 245 250 255
 Gln Leu Ser Pro Leu Glu Met Glu Leu Ala Asp Arg Tyr Leu Ala Gly
 260 265 270
 Gln Thr Glu Asp Asn Asp Ser Gly Ser Glu His Ser Lys Pro Ser Asp
 275 280 285
 Lys Glu Val Thr His Thr Phe Leu Gly His Arg Ile Lys Ala Tyr Gly
 290 295 300
 Lys Gly Leu Asp Gly Lys Pro Tyr Asp Thr Ser Asp Ala Tyr Val Phe
 305 310 315 320
 Ser Lys Glu Ser Ile His Ser Val Asp Lys Ser Gly Val Thr Ala Lys
 325 330 335
 His Gly Asp His Phe His Tyr Ile Gly Phe Gly Glu Leu Glu Gln Tyr
 340 345 350

Glu Leu Asp Glu Val Ala Asn Trp Val Lys Ala Lys Gly Gln Ala Asp
 355 360 365
 Glu Leu Ala Ala Ala Leu Asp Gln Glu Gln Gly Lys Glu Lys Pro Leu
 370 375 380
 Phe Asp Thr Lys Lys Val Ser Arg Lys Val Thr Lys Asp Gly Lys Val
 385 390 395 400
 Gly Tyr Met Met Pro Lys Asp Gly Lys Asp Tyr Phe Tyr Ala Arg Asp
 405 410 415
 Gln Leu Asp Leu Thr Gln Ile Ala Phe Ala Glu Gln Glu Leu Met Leu
 420 425 430
 Lys Asp Lys Lys His Tyr Arg Tyr Asp Ile Val Asp Thr Gly Ile Glu
 435 440 445
 Pro Arg Leu Ala Val Asp Val Ser Ser Leu Pro Met His Ala Gly Asn
 450 455 460
 Ala Thr Tyr Asp Thr Gly Ser Ser Phe Val Ile Pro His Ile Asp His
 465 470 475 480
 Ile His Val Val Pro Tyr Ser Trp Leu Thr Arg Asp Gln Ile Ala Thr
 485 490 495
 Val Lys Tyr Val Met Gln His Pro Glu Val Arg Pro Asp Val Trp Ser
 500 505 510
 Lys Pro Gly His Glu Glu Ser Gly Ser Val Ile Pro Asn Val Thr Pro
 515 520 525
 Leu Asp Lys Arg Ala Gly Met Pro Asn Trp Gln Ile Ile His Ser Ala
 530 535 540
 Glu Glu Val Gln Lys Ala Leu Ala Glu Gly Arg Phe Ala Thr Pro Asp
 545 550 555 560
 Gly Tyr Ile Phe Asp Pro Arg Asp Val Leu Ala Lys Glu Thr Phe Val
 565 570 575
 Trp Lys Asp Gly Ser Phe Ser Ile Pro Arg Ala Asp Gly Ser Ser Leu
 580 585 590
 Arg Thr Ile Asn Lys Ser Asp Leu Ser Gln Ala Glu Trp Gln Gln Ala
 595 600 605
 Gln Glu Leu Leu Ala Lys Lys Asn Thr Gly Asp Ala Thr Asp Thr Asp
 610 615 620
 Lys Pro Lys Glu Lys Gln Gln Ala Asp Lys Ser Asn Glu Asn Gln Gln
 625 630 635 640
 Pro Ser Glu Ala Ser Lys Glu Glu Lys Glu Ser Asp Asp Phe Ile Asp
 645 650 655
 Ser Leu Pro Asp Tyr Gly Leu Asp Arg Ala Thr Leu Glu Asp His Ile
 660 665 670
 Asn Gln Leu Ala Gln Lys Ala Asn Ile Asp Pro Lys Tyr Leu Ile Phe
 675 680 685
 Gln Pro Glu Gly Val Gln Phe Tyr Asn Lys Asn Gly Glu Leu Val Thr
 690 695 700
 Tyr Asp Ile Lys Thr Leu Gln Gln Ile Asn Pro
 705 710 715

<210> 17
 <211> 77
 <212> PRT
 <213> Streptococcus

<400> 17

Met His Ser Phe Ser Asn Pro Gly Tyr Pro Tyr Asp Asn Ala Val Thr
 1 5 10 15

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Glu Ala Phe Phe Lys Tyr Leu Lys His Arg Gln Ile Asn Arg Lys His
 20 25 30
 Tyr Gln Asn Ile Lys Gln Val Gln Leu Asp Cys Phe Glu Tyr Ile Glu
 35 40 45
 Asn Phe Tyr Asn Asn Tyr Asn Pro His Thr Ala Asn Leu Gly Leu Thr
 50 55 60
 Pro Asn Gln Lys Glu Glu Asn Tyr Phe Asn Ala Ile Lys
 65 70 75

<210> 18
 <211> 86
 <212> PRT
 <213> Streptococcus

<400> 18
 Met Ala Tyr Tyr Gln Ala Cys Thr Glu Lys Asp Ile Ile Arg Ser Met
 1 5 10 15
 Ser Arg Lys Gly Thr Pro Ala Asp Asn Ala Cys Ile Glu Trp Phe His
 20 25 30
 Thr Val Leu Lys Thr Glu Thr Phe Tyr Phe His Asn Arg Arg Lys Tyr
 35 40 45
 Asn Lys Asp Ser Ile Thr Asn Ile Val Lys Asn Tyr Ile Thr Phe Tyr
 50 55 60
 Asn Glu Thr Arg Ile Gln Gln Arg Leu Asn Asp Gln Ser Pro Val Gln
 65 70 75 80
 Tyr Arg Lys Leu Ile Ala
 85

<210> 19
 <211> 126
 <212> PRT
 <213> Streptococcus

<400> 19
 Met Glu Asn His Phe Ile Tyr Gly Tyr Arg Thr Ile Thr Arg Leu Leu
 1 5 10 15
 Lys Lys Ile His Gly Leu Thr Val Asn Thr Lys Lys Val Tyr Arg Ile
 20 25 30
 Met Lys Asn Asn Gly Trp Leu Cys Arg Thr Arg Thr Lys Lys Val Pro
 35 40 45
 Asn Leu Gly Lys Ala Tyr Tyr Leu Thr Asp Asn Lys Leu Ser Arg Asp
 50 55 60
 Phe His Ala Asp Lys Pro Lys Glu Lys Leu Val Thr Asp Ile Thr Tyr
 65 70 75 80
 Leu Tyr Phe Gly Asn Cys Lys Leu Tyr Leu Ser Ser Ile Met Asn Leu
 85 90 95
 Tyr Asn Arg Glu Ile Ile Ala Tyr Thr Ile Ser Asp Cys Gln Asp Thr
 100 105 110
 Asp Phe Val Leu Asp Thr Leu Asn Gln Leu Lys Leu Pro Lys
 115 120 125

<210> 20
 <211> 96
 <212> PRT
 <213> Streptococcus

<400> 20
 Met Val Lys Lys Ala Tyr Ser Trp Glu Thr Lys Leu Ala Cys Ile Asp
 1 5 10 15
 Met Lys Lys Ala Gly Lys Ser Asn Arg Val Ile Met Glu Thr Leu Gly
 20 25 30
 Ile Lys Asn Asn Ser Gln Ile Tyr Thr Trp Met Lys Trp Tyr Glu Asn
 35 40 45
 Glu Glu Leu Tyr Arg Phe His Gln Gly Val Gly Lys Gln Tyr Thr Tyr
 50 55 60
 Gly Lys Gly Leu Glu His Leu Ser Glu Val Glu Gln Leu Gln Leu Gln
 65 70 75 80
 Val Asp Leu Leu Lys Tyr Arg Gly Leu Ile Arg Lys Ser Ile Lys
 85 90 95

<210> 21
 <211> 288
 <212> PRT
 <213> streptococcus

<400> 21
 Ile Arg Tyr Pro Lys Ala Ser Ser Gly Asp Tyr Gly Thr Lys Arg Glu
 1 5 10 15
 Ile Ile Thr Ala Asn Lys Asp Lys Tyr Ser Ile Ser Lys Met Cys Arg
 20 25 30
 Trp Leu Asn Met Pro His Ser Ser Tyr Tyr Tyr Gln Ala Val Glu Ser
 35 40 45
 Val Ser Glu Thr Glu Phe Glu Glu Thr Ile Lys Arg Ile Phe Leu Asp
 50 55 60
 Ser Glu Ser Arg Tyr Gly Ser Arg Lys Ile Lys Ile Cys Leu Asn Asn
 65 70 75 80
 Glu Gly Ile Thr Leu Ser Arg Arg Arg Ile Arg Arg Ile Met Lys Arg
 85 90 95
 Leu Asn Leu Val Ser Val Tyr Gln Lys Ala Thr Phe Lys Pro His Ser
 100 105 110
 Arg Gly Lys Asn Glu Ala Pro Ile Pro Asn His Leu Asp Arg Gln Phe
 115 120 125
 Lys Gln Glu Arg Pro Leu Gln Ala Leu Val Thr Asp Leu Thr Tyr Val
 130 135 140
 Arg Val Gly Asn Arg Trp Ala Tyr Val Cys Leu Ile Ile Asp Leu Tyr
 145 150 155 160
 Asn Arg Glu Ile Ile Gly Leu Ser Leu Gly Trp His Lys Thr Ala Glu
 165 170 175
 Leu Val Lys Gln Ala Ile Gln Ser Ile Pro Tyr Ala Leu Thr Lys Val
 180 185 190
 Lys Met Phe His Ser Asp Arg Gly Lys Glu Phe Asp Asn Gln Leu Ile
 195 200 205
 Asp Glu Ile Leu Glu Ala Phe Gly Ile Thr Arg Ser Leu Ser Gln Ala
 210 215 220
 Gly Tyr Pro Tyr Asp Asn Ala Val Ala Glu Ser Thr Tyr Arg Ala Phe
 225 230 235 240
 Lys Ile Glu Phe Val Tyr Gln Glu Thr Phe Gln Leu Leu Glu Glu Leu
 245 250 255
 Ala Leu Lys Thr Lys Asp Tyr Val His Trp Trp Asn Tyr His Arg Ile
 260 265 270
 His Gly Ser Leu Asn Tyr Gln Thr Pro Met Thr Lys Arg Leu Ile Ala
 275 280 285

<210> 22
 <211> 5058
 <212> DNA
 <213> streptococcus

<220>
 <221> CDS
 <222> (1) ... (663)

<221> CDS
 <222> (763) ... (1344)

<221> CDS
 <222> (1362) ... (1739)

<221> CDS
 <222> (2266) ... (5058)

<400> 22

aat ttg aaa gca gaa tta tct gta gaa gat gag caa tat aca gca aca	48
Asn Leu Lys Ala Glu Leu Ser Val Glu Asp Glu Gln Tyr Thr Ala Thr	
1 5 10 15	
gtt tat ggt aaa tct gct cat ggt tca aca cca caa gaa ggt gtt aat	96
Val Tyr Gly Lys Ser Ala His Gly Ser Thr Pro Gln Glu Gly Val Asn	
20 25 30	
ggg gcg act tat tta gct ctt tat cta agt caa ttt gat ttt gaa ggt	144
Gly Ala Thr Tyr Leu Ala Leu Tyr Leu Ser Gln Phe Asp Phe Glu Gly	
35 40 45	
cct gct cgt gct ttc tta gat gtt aca gcc aac att att cac gaa gac	192
Pro Ala Arg Ala Phe Leu Asp Val Thr Ala Asn Ile Ile His Glu Asp	
50 55 60	
ttc tca ggt gaa aaa ctt gga gta gct tat gaa gat gac tgt atg gga	240
Phe Ser Gly Glu Lys Leu Gly Val Ala Tyr Glu Asp Asp Cys Met Gly	
65 70 75 80	
cca ttg agc atg aat gca ggt gtc ttc cag ttt gat gaa act aat gat	288
Pro Leu Ser Met Asn Ala Gly Val Phe Gln Phe Asp Glu Thr Asn Asp	
85 90 95	
gat aat act atc gct ctt aat ttc cgt tac cca caa ggg aca gat gct	336
Asp Asn Thr Ile Ala Leu Asn Phe Arg Tyr Pro Gln Gly Thr Asp Ala	
100 105 110	
aaa act atc caa act aag ctt gag aaa ctt aac gga gtt gaa aaa gtg	384
Lys Thr Ile Gln Thr Lys Leu Glu Lys Leu Asn Gly Val Glu Lys Val	
115 120 125	
act ctt tct gac cat gaa cac aca cca cac tat gta cct atg gac gat	432
Thr Leu Ser Asp His Glu His Thr Pro His Tyr Val Pro Met Asp Asp	
130 135 140	

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gaa tta gta tca acc tta cta gct gtc tat gaa aag caa act ggt ctt	480
Glu Leu Val Ser Thr Leu Leu Ala Val Tyr Glu Lys Gln Thr Gly Leu	
145 150 155 160	
aaa gga cat gaa cag gtt att ggt ggt ggg aca ttt ggt cgc tta ctt	528
Lys Gly His Glu Gln Val Ile Gly Gly Thr Phe Gly Arg Leu Leu	
165 170 175	
gaa cgg ggt gtt gca tac ggt gcc atg ttc cca gga gat gaa aac act	576
Glu Arg Gly Val Ala Tyr Gly Ala Met Phe Pro Gly Asp Glu Asn Thr	
180 185 190	
atg cat caa gct aat gag tac atg cct tta gaa aat att ttc cgt tcg	624
Met His Gln Ala Asn Glu Tyr Met Pro Leu Glu Asn Ile Phe Arg Ser	
195 200 205	
gct gct atc tac gca gaa gct atc tat gaa tta atc aaa taaaataatc	673
Ala Ala Ile Tyr Ala Glu Ala Ile Tyr Glu Leu Ile Lys	
210 215 220	
cttaaactaa atatgtgatc aatgataaaag ggtggtaag acatgaaagt gtcttgccct	733
cttttcataa ggttagattt ggagacttt atg act gac ttg gaa aaa att att	786
Met Thr Asp Leu Glu Lys Ile Ile	
225	
aaa gca ata aaa agt gat tca cag aat caa aat tat aca gaa aat ggt	834
Lys Ala Ile Lys Ser Asp Ser Gln Asn Gln Asn Tyr Thr Glu Asn Gly	
230 235 240 245	
att gat cct ttg ttt gct gct cct aaa aca gct agg atc aat att gtt	882
Ile Asp Pro Leu Phe Ala Ala Pro Lys Thr Ala Arg Ile Asn Ile Val	
250 255 260	
ggc caa gca cct ggt tta aaa act caa gaa gca aga ctc tat tgg aaa	930
Gly Gln Ala Pro Gly Leu Lys Thr Gln Glu Ala Arg Leu Tyr Trp Lys	
265 270 275	
gat aaa tct gga gat cgt cta cgc cag tgg ctt gga gtt gat gaa gag	978
Asp Lys Ser Gly Asp Arg Leu Arg Gln Trp Leu Gly Val Asp Glu Glu	
280 285 290	
aca ttt tac cat tct gga aaa ttt gct gtt tta cct tta gat ttt tat	1026
Thr Phe Tyr His Ser Gly Lys Phe Ala Val Leu Pro Leu Asp Phe Tyr	
295 300 305	
tac cca ggc aaa gga aaa tca gga gat tta ccc cct aga aaa ggt ttt	1074
Tyr Pro Gly Lys Gly Lys Ser Gly Asp Leu Pro Pro Arg Lys Gly Phe	
310 315 320 325	
gcg gag aaa tgg cac cct ctt att tta aaa gaa atg cct aat gtt caa	1122
Ala Glu Lys Trp His Pro Leu Ile Leu Lys Glu Met Pro Asn Val Gln	
330 335 340	
ttg acc ttg cta gtt ggt cag tat gct cag aaa tat tat ctt gga agc	1170
Leu Thr Leu Leu Val Gly Gln Tyr Ala Gln Lys Tyr Tyr Leu Gly Ser	
345 350 355	

tcc gca cat aaa aat cta aca gaa aca gtt aaa gct tac aaa gac tat Ser Ala His Lys Asn Leu Thr Glu Thr Val Lys Ala Tyr Lys Asp Tyr 360 365 370	1218
ctt ccc gat tat tta ccc ctg gtt cac cca tca ccg cga aat caa att Leu Pro Asp Tyr Leu Pro Leu Val His Pro Ser Pro Arg Asn Gln Ile 375 380 385	1266
tgg cta aag aag aat cca tgg ttt gaa aaa gat cta atc gtt gat tta Trp Leu Lys Lys Asn Pro Trp Phe Glu Lys Asp Leu Ile Val Asp Leu 390 395 400 405	1314
caa aag ata gta gca gat att tta aaa gat taaggatagg agttgggt atg Gln Lys Ile Val Ala Asp Ile Leu Lys Asp Met 410 415	1364
aga gat aat cat cta cac acg tat ttt tcc tat gat tgt caa acg gca Arg Asp Asn His Leu His Thr Tyr Phe Ser Tyr Asp Cys Gln Thr Ala 420 425 430	1412
ttt gag gac tat att aat ggt ttt aca ggt gaa ttt atc acg aca gaa Phe Glu Asp Tyr Ile Asn Gly Phe Thr Gly Glu Phe Ile Thr Thr Glu 435 440 445	1460
cat ttt gat tta tca aat cct tac acc ggt caa gac gat gtt cct gat His Phe Asp Leu Ser Asn Pro Tyr Thr Gly Gln Asp Asp Val Pro Asp 450 455 460	1508
tat agt gct tat tgt caa aaa ata gat tat ctt aat cag aaa tat gga Tyr Ser Ala Tyr Cys Gln Lys Ile Asp Tyr Leu Asn Gln Lys Tyr Gly 465 470 475 480	1556
aat cga ttt aaa aaa gga att gaa atc ggt tat ttt aaa gat agg gaa Asn Arg Phe Lys Lys Gly Ile Glu Ile Gly Tyr Phe Lys Asp Arg Glu 485 490 495	1604
tca gat att tta gat tat tta aaa aat aaa gaa ttt gat tta aaa cta Ser Asp Ile Leu Asp Tyr Leu Lys Asn Lys Glu Phe Asp Leu Lys Leu 500 505 510	1652
ttg tca atc cat cat aat ggt agg tat gat tat ctg caa gaa gaa gct Leu Ser Ile His His Asn Gly Arg Tyr Asp Tyr Leu Gln Glu Glu Ala 515 520 525	1700
ctg aaa gta cca aca aag gga gct ttt agc aga tta ctt taatcgatg Leu Lys Val Pro Thr Lys Gly Ala Phe Ser Arg Leu Leu 530 535 540	1749
gaatttgcataggccgtgt ggaagcgcac gtttagctc actttgatta tggtttcgt aagtttaact tagatgtaga agatttaaaa ccgttgaaa cgcaattgaa ggcgcattttc ataaaagatgt tatctaagggtt gtagctttt gaaactaata ccaaatccct ttatctata ggaaatggaaa aactttatcg ctatgtttt gagataactca aacagcttgg ttgtaaacaa tactctatag gctctgacgg tcatattcct gaacatttt gttatgaatt tgatagactt caaggctctgc taaaggacta tcaaattgat gaaaatcatt tgatatgagg aaattttga taaaaaaagct aggcaatatt gcttagctt tttgtatgc tattgtatgt tttgtatgaaa	1809 1869 1929 1989 2049 2109 2169

tttcaaaaa aataaaagaaa tcatttactt gttgcaagcg cttgcgtaaa ttgttatgt	2229
tttattggta acaattcatt aaaaaaggag aatgat atg aaa aga aaa gac tta	2283
Met Lys Arg Lys Asp Leu	
545	
ttt ggt gat aaa caa act caa tac acg att aga aag tta agt gtt gga	2331
Phe Gly Asp Lys Gln Thr Gln Tyr Thr Ile Arg Lys Leu Ser Val Gly	
550 555 560	
gta gct tca gtt aca aca ggg gta tgt att ttt ctt cat agt cca cag	2379
Val Ala Ser Val Thr Thr Gly Val Cys Ile Phe Leu His Ser Pro Gln	
565 570 575	
gta ttt gct gaa gaa gta agt gtt tct cct gca act aca gcg att gca	2427
Val Phe Ala Glu Glu Val Ser Val Ser Pro Ala Thr Thr Ala Ile Ala	
580 585 590 595	
gag tcg aat att aat cag gtt gac aac caa caa tct act aat tta aaa	2475
Glu Ser Asn Ile Asn Gln Val Asp Asn Gln Gln Ser Thr Asn Leu Lys	
600 605 610	
gat gac ata aac tca aac tct gag acg gtt gtg aca ccc tca gat atg	2523
Asp Asp Ile Asn Ser Asn Ser Glu Thr Val Val Thr Pro Ser Asp Met	
615 620 625	
ccg gat acc aag caa tta gta tca gat gaa act gac act caa aag gga	2571
Pro Asp Thr Lys Gln Leu Val Ser Asp Glu Thr Asp Thr Gln Lys Gly	
630 635 640	
gtg aca gag ccg gat aag gcg aca agc ctg ctt gaa gaa aat aaa ggt	2619
Val Thr Glu Pro Asp Lys Ala Thr Ser Leu Leu Glu Glu Asn Lys Gly	
645 650 655	
cct gtt tca gat aaa aat acc tta gat tta aaa gta gca cca tct aca	2667
Pro Val Ser Asp Lys Asn Thr Leu Asp Leu Lys Val Ala Pro Ser Thr	
660 665 670 675	
ttg caa aat act ccc gac aaa act tct caa gct ata ggt gct cca agc	2715
Leu Gln Asn Thr Pro Asp Lys Thr Ser Gln Ala Ile Gly Ala Pro Ser	
680 685 690	
cct acc ttg aaa gta gct aat caa gct cca cgg att gaa aat ggt tac	2763
Pro Thr Leu Lys Val Ala Asn Gln Ala Pro Arg Ile Glu Asn Gly Tyr	
695 700 705	
ttt agg cta cat ctt aaa gaa ttg cct caa ggt cat cct gta gaa agc	2811
Phe Arg Leu His Leu Lys Glu Leu Pro Gln Gly His Pro Val Glu Ser	
710 715 720	
act gga ctt tgg ata tgg gga gat gtt gat caa ccc tct agt aat tgg	2859
Thr Gly Leu Trp Ile Trp Gly Asp Val Asp Gln Pro Ser Ser Asn Trp	
725 730 735	
cca aat ggt gct atc cct atg act gat gct aag aaa gat gat tac ggt	2907
Pro Asn Gly Ala Ile Pro Met Thr Asp Ala Lys Lys Asp Asp Tyr Gly	
740 745 750 755	

tat tat gtt gat ttt aaa tta tct gaa aaa caa cga aaa caa ata tct Tyr Tyr Val Asp Phe Lys Leu Ser Glu Lys Gln Arg Lys Gln Ile Ser	2955
760 765 770	
ttt tta att aat aac aaa gca ggg aca aat tta agc ggc gat cat cat Phe Leu Ile Asn Asn Lys Ala Gly Thr Asn Leu Ser Gly Asp His His	3003
775 780 785	
att cca tta tta cga cct gag atg aac caa gtt tgg att gat gaa aag Ile Pro Leu Leu Arg Pro Glu Met Asn Gln Val Trp Ile Asp Glu Lys	3051
790 795 800	
tac ggt ata cat act tat caa ccc ctc aaa gaa ggg tat gtc cgt att Tyr Gly Ile His Thr Tyr Gln Pro Leu Lys Glu Gly Tyr Val Arg Ile	3099
805 810 815	
aac tat ttg agt tcc tct agt aac tat gac cac tta tca gca tgg ctc Asn Tyr Leu Ser Ser Asn Tyr Asp His Leu Ser Ala Trp Leu	3147
820 825 830 835	
ttt aaa gat gtt gca acc ccy tca aca act tgg cca gat ggt agt aat Phe Lys Asp Val Ala Thr Xaa Ser Thr Thr Trp Pro Asp Gly Ser Asn	3195
840 845 850	
ttt gtg aat caa gga cta tat gga agg tat att gat gta tca cta aaa Phe Val Asn Gln Gly Leu Tyr Gly Arg Tyr Ile Asp Val Ser Leu Lys	3243
855 860 865	
act aac gcc aaa gag att ggt ttt cta atc tta gat gaa agt aag aca Thr Asn Ala Lys Glu Ile Gly Phe Leu Ile Leu Asp Glu Ser Lys Thr	3291
870 875 880	
gga gat gca gtg aaa gtt caa ccc aac gac tat gtt ttt aga gat tta Gly Asp Ala Val Lys Val Gln Pro Asn Asp Tyr Val Phe Arg Asp Leu	3339
885 890 895	
gct aac cat aac caa att ttt gta aaa gat aag gat cca aag gtt tat Ala Asn His Asn Gln Ile Phe Val Lys Asp Lys Asp Pro Lys Val Tyr	3387
900 905 910 915	
aat aat cct tat tac att gat caa gtg cag cta aag gat gcc caa caa Asn Asn Pro Tyr Tyr Ile Asp Gln Val Gln Leu Lys Asp Ala Gln Gln	3435
920 925 930	
att gat tta aca agt att caa gca agt ttt aca act cta gat ggg gta Ile Asp Leu Thr Ser Ile Gln Ala Ser Phe Thr Thr Leu Asp Gly Val	3483
935 940 945	
gat aaa act gaa att tta aaa gaa ttg aaa gtg act gat aaa aat caa Asp Lys Thr Glu Ile Leu Lys Glu Leu Lys Val Thr Asp Lys Asn Gln	3531
950 955 960	
aat gct ata caa att tct gat atc act ctc gat act agt aaa tct ctt Asn Ala Ile Gln Ile Ser Asp Ile Thr Leu Asp Thr Ser Lys Ser Leu	3579
965 970 975	

tta ata atc aaa ggc gac ttt aat cct aaa caa ggt cat ttc aac ata	3627
Leu Ile Ile Lys Gly Asp Phe Asn Pro Lys Gln Gly His Phe Asn Ile	
980 985 990 995	
tct tat aat ggt aac aat gtc atg aca agg caa tct tgg gaa ttt aaa	3675
Ser Tyr Asn Gly Asn Asn Val Met Thr Arg Gln Ser Trp Glu Phe Lys	
1000 1005 1010	
gac caa ctt tat gct tat agt gga aat tta ggt gca gtt ctc aat caa	3723
Asp Gln Leu Tyr Ala Tyr Ser Gly Asn Leu Gly Ala Val Leu Asn Gln	
1015 1020 1025	
gat ggt tca aaa gtt gaa gcc agc ctc tgg tca ccg agt gct gat agt	3771
Asp Gly Ser Lys Val Glu Ala Ser Leu Trp Ser Pro Ser Ala Asp Ser	
1030 1035 1040 1045	
gtc act atg att att tat gac aaa gat aac caa aac agg gtt gta gcg	3819
Val Thr Met Ile Ile Tyr Asp Lys Asp Asn Gln Asn Arg Val Val Ala	
1045 1050 1055	
act acc ccc ctt gtg aaa aat aat aaa ggt gtt tgg cag acg ata ctt	3867
Thr Thr Pro Leu Val Lys Asn Asn Lys Gly Val Trp Gln Thr Ile Leu	
1060 1065 1070 1075	
gat act aaa tta ggt att aaa aac tat act ggt tac tat tat ctt tac	3915
Asp Thr Lys Leu Gly Ile Lys Asn Tyr Thr Gly Tyr Tyr Tyr Leu Tyr	
1080 1085 1090	
gaa ata aaa aga ggt aag gat aag gtt aag att tta gat cct tat gca	3963
Glu Ile Lys Arg Gly Lys Asp Lys Val Lys Ile Leu Asp Pro Tyr Ala	
1095 1100 1105	
aag tca tta gca gag tgg gat agt aat act gtt aat gat gat att aaa	4011
Lys Ser Leu Ala Glu Trp Asp Ser Asn Thr Val Asn Asp Asp Ile Lys	
1110 1115 1120	
acg gct aaa gca gct ttt gta aat cca agt caa ctt gga cct caa aat	4059
Thr Ala Lys Ala Ala Phe Val Asn Pro Ser Gln Leu Gly Pro Gln Asn	
1125 1130 1135	
tta agt ttt gct aaa att gct aat ttt aaa gga aga caa gat gct gtt	4107
Leu Ser Phe Ala Lys Ile Ala Asn Phe Lys Gly Arg Gln Asp Ala Val	
1140 1145 1150 1155	
ata tac gaa gca cat gta aga gac ttc act tct gat cga tct ttg gat	4155
Ile Tyr Glu Ala His Val Arg Asp Phe Thr Ser Asp Arg Ser Leu Asp	
1160 1165 1170	
gga aaa tta aaa aat caa ttt ggt acc ttt gca gcc ttt tca gag aaa	4203
Gly Lys Leu Lys Asn Gln Phe Gly Thr Phe Ala Ala Phe Ser Glu Lys	
1175 1180 1185	
cta gat tat tta cag aaa tta gga gtt aca cac att cag ctt tta ccg	4251
Leu Asp Tyr Leu Gln Lys Leu Gly Val Thr His Ile Gln Leu Leu Pro	
1190 1195 1200	

gta ttg agt tat ttt tat gtt aat gaa atg gat aag tca cgc tca aca	4299
Val Leu Ser Tyr Phe Tyr Val Asn Glu Met Asp Lys Ser Arg Ser Thr	
1205 1210 1215	
gct tac act tcc tca gac aat aat tac aat tgg ggc tat gac cca cag	4347
Ala Tyr Thr Ser Ser Asp Asn Asn Tyr Asn Trp Gly Tyr Asp Pro Gln	
1220 1225 1230 1235	
agc tat ttt gct ctt tct ggg atg tat tca gag aaa cca aaa gat cca	4395
Ser Tyr Phe Ala Leu Ser Gly Met Tyr Ser Glu Lys Pro Lys Asp Pro	
1240 1245 1250	
tca gca cgt atc gcc gaa tta aaa caa tta ata cat gat att cat aaa	4443
Ser Ala Arg Ile Ala Glu Leu Lys Gln Leu Ile His Asp Ile His Lys	
1255 1260 1265	
cgt ggc atg ggg gtt ata ctt gat gtc gtc tat aat cac act gca aaa	4491
Arg Gly Met Gly Val Ile Leu Asp Val Val Tyr Asn His Thr Ala Lys	
1270 1275 1280	
act tat ctc ttt gag gat ata gaa cct aat tat tat cac ttt atg aat	4539
Thr Tyr Leu Phe Glu Asp Ile Glu Pro Asn Tyr Tyr His Phe Met Asn	
1285 1290 1295	
gaa gat ggt tca cca aga gaa agt ttt gga ggg gga cgt tta gga acc	4587
Glu Asp Gly Ser Pro Arg Glu Ser Phe Gly Gly Arg Leu Gly Thr	
1300 1305 1310 1315	
act cat gca atg agt cgt cgt gtt ttg gtt gat tcc att aaa tat ctt	4635
Thr His Ala Met Ser Arg Arg Val Leu Val Asp Ser Ile Lys Tyr Leu	
1320 1325 1330	
aca agt gaa ttt aaa gtt gat ggt ttc cgt ttt gat atg atg gga gat	4683
Thr Ser Glu Phe Lys Val Asp Gly Phe Arg Phe Asp Met Met Gly Asp	
1335 1340 1345	
cat gat gcg gct gcg att gaa tta gct tat aaa gaa gct aaa gct att	4731
His Asp Ala Ala Ala Ile Glu Leu Ala Tyr Lys Glu Ala Lys Ala Ile	
1350 1355 1360	
aat cct aat atg att atg att ggt gag ggc tgg aga aca ttc caa ggc	4779
Asn Pro Asn Met Ile Met Ile Gly Glu Gly Trp Arg Thr Phe Gln Gly	
1365 1370 1375	
gat caa ggt cag ccg gtt aaa cca gct gac caa gat tgg atg aag tca	4827
Asp Gln Gly Gln Pro Val Lys Pro Ala Asp Gln Asp Trp Met Lys Ser	
1380 1385 1390 1395	
acc gat aca gtt ggc gtc ttt tca gat gat att cgt aat agc ttg aaa	4875
Thr Asp Thr Val Gly Val Phe Ser Asp Asp Ile Arg Asn Ser Leu Lys	
1400 1405 1410	
tct ggt ttt cca aat gaa ggt act cca gct ttc atc aca ggt ggc cca	4923
Ser Gly Phe Pro Asn Glu Gly Thr Pro Ala Phe Ile Thr Gly Gly Pro	
1415 1420 1425	

caa tct tta caa ggt att ttt aaa aat atc aaa gca caa cct ggg aat 4971
 Gln Ser Leu Gln Gly Ile Phe Lys Asn Ile Lys Ala Gln Pro Gly Asn
 1430 1435 1440

ttt gaa gca gat tcg cca gga gat gtg gtg cag tat att gct gca cat 5019
 Phe Glu Ala Asp Ser Pro Gly Asp Val Val Gln Tyr Ile Ala Ala His
 1445 1450 1455

gat aac ctt acc ttg cat gat gtg att gca aaa tca att 5058
 Asp Asn Leu Thr Leu His Asp Val Ile Ala Lys Ser Ile
 1460 1465 1470

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 35 40 45
 Pro Ala Arg Ala Phe Leu Asp Val Thr Ala Asn Ile Ile His Glu Asp
 50 55 60
 Phe Ser Gly Glu Lys Leu Gly Val Ala Tyr Glu Asp Asp Cys Met Gly
 65 70 75 80
 Pro Leu Ser Met Asn Ala Gly Val Phe Gln Phe Asp Glu Thr Asn Asp
 85 90 95
 Asp Asn Thr Ile Ala Leu Asn Phe Arg Tyr Pro Gln Gly Thr Asp Ala
 100 105 110
 Lys Thr Ile Gln Thr Lys Leu Glu Lys Leu Asn Gly Val Glu Lys Val
 115 120 125
 Thr Leu Ser Asp His Glu His Thr Pro His Tyr Val Pro Met Asp Asp
 130 135 140
 Glu Leu Val Ser Thr Leu Leu Ala Val Tyr Glu Lys Gln Thr Gly Leu
 145 150 155 160
 Lys Gly His Glu Gln Val Ile Gly Gly Thr Phe Gly Arg Leu Leu
 165 170 175
 Glu Arg Gly Val Ala Tyr Gly Ala Met Phe Pro Gly Asp Glu Asn Thr
 180 185 190
 Met His Gln Ala Asn Glu Tyr Met Pro Leu Glu Asn Ile Phe Arg Ser
 195 200 205
 Ala Ala Ile Tyr Ala Glu Ala Ile Tyr Glu Leu Ile Lys
 210 215 220

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<400> 24

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 Lys Thr Ala Arg Ile Asn Ile Val Gly Gln Ala Pro Gly Leu Lys Thr
 35 40 45
 Gln Glu Ala Arg Leu Tyr Trp Lys Asp Lys Ser Gly Asp Arg Leu Arg
 50 55 60
 Gln Trp Leu Gly Val Asp Glu Glu Thr Phe Tyr His Ser Gly Lys Phe
 65 70 75 80
 Ala Val Leu Pro Leu Asp Phe Tyr Tyr Pro Gly Lys Gly Lys Ser Gly
 85 90 95
 Asp Leu Pro Pro Arg Lys Gly Phe Ala Glu Lys Trp His Pro Leu Ile
 100 105 110
 Leu Lys Glu Met Pro Asn Val Gln Leu Thr Leu Leu Val Gly Gln Tyr
 115 120 125
 Ala Gln Lys Tyr Tyr Leu Gly Ser Ser Ala His Lys Asn Leu Thr Glu
 130 135 140
 Thr Val Lys Ala Tyr Lys Asp Tyr Leu Pro Asp Tyr Leu Pro Leu Val
 145 150 155 160
 His Pro Ser Pro Arg Asn Gln Ile Trp Leu Lys Lys Asn Pro Trp Phe
 165 170 175
 Glu Lys Asp Leu Ile Val Asp Leu Gln Lys Ile Val Ala Asp Ile Leu
 180 185 190
 Lys Asp

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 Ala Phe Glu Asp Tyr Ile Asn Gly Phe Thr Gly Glu Phe Ile Thr Thr
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 35 40 45
 Asp Tyr Ser Ala Tyr Cys Gln Lys Ile Asp Tyr Leu Asn Gln Lys Tyr
 50 55 60
 Gly Asn Arg Phe Lys Lys Gly Ile Glu Ile Gly Tyr Phe Lys Asp Arg
 65 70 75 80
 Glu Ser Asp Ile Leu Asp Tyr Leu Lys Asn Lys Glu Phe Asp Leu Lys
 85 90 95
 Leu Leu Ser Ile His His Asn Gly Arg Tyr Asp Tyr Leu Gln Glu Glu
 100 105 110
 Ala Leu Lys Val Pro Thr Lys Gly Ala Phe Ser Arg Leu Leu
 115 120 125

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<400> 26

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 Arg Lys Leu Ser Val Gly Val Ala Ser Val Thr Thr Gly Val Cys Ile
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 Phe Leu His Ser Pro Gln Val Phe Ala Glu Glu Val Ser Val Ser Pro
 35 40 45
 Ala Thr Thr Ala Ile Ala Glu Ser Asn Ile Asn Gln Val Asp Asn Gln
 50 55 60
 Gln Ser Thr Asn Leu Lys Asp Asp Ile Asn Ser Asn Ser Glu Thr Val
 65 70 75 80
 Val Thr Pro Ser Asp Met Pro Asp Thr Lys Gln Leu Val Ser Asp Glu
 85 90 95
 Thr Asp Thr Gln Lys Gly Val Thr Glu Pro Asp Lys Ala Thr Ser Leu
 100 105 110
 Leu Glu Glu Asn Lys Gly Pro Val Ser Asp Lys Asn Thr Leu Asp Leu
 115 120 125
 Lys Val Ala Pro Ser Thr Leu Gln Asn Thr Pro Asp Lys Thr Ser Gln
 130 135 140
 Ala Ile Gly Ala Pro Ser Pro Thr Leu Lys Val Ala Asn Gln Ala Pro
 145 150 155 160
 Arg Ile Glu Asn Gly Tyr Phe Arg Leu His Leu Lys Glu Leu Pro Gln
 165 170 175
 Gly His Pro Val Glu Ser Thr Gly Leu Trp Ile Trp Gly Asp Val Asp
 180 185 190
 Gln Pro Ser Ser Asn Trp Pro Asn Gly Ala Ile Pro Met Thr Asp Ala
 195 200 205
 Lys Lys Asp Asp Tyr Gly Tyr Tyr Val Asp Phe Lys Leu Ser Glu Lys
 210 215 220
 Gln Arg Lys Gln Ile Ser Phe Leu Ile Asn Asn Lys Ala Gly Thr Asn
 225 230 235 240
 Leu Ser Gly Asp His His Ile Pro Leu Leu Arg Pro Glu Met Asn Gln
 245 250 255
 Val Trp Ile Asp Glu Lys Tyr Gly Ile His Thr Tyr Gln Pro Leu Lys
 260 265 270
 Glu Gly Tyr Val Arg Ile Asn Tyr Leu Ser Ser Ser Asn Tyr Asp
 275 280 285
 His Leu Ser Ala Trp Leu Phe Lys Asp Val Ala Thr Xaa Ser Thr Thr
 290 295 300
 Trp Pro Asp Gly Ser Asn Phe Val Asn Gln Gly Leu Tyr Gly Arg Tyr
 305 310 315 320
 Ile Asp Val Ser Leu Lys Thr Asn Ala Lys Glu Ile Gly Phe Leu Ile
 325 330 335
 Leu Asp Glu Ser Lys Thr Gly Asp Ala Val Lys Val Gln Pro Asn Asp
 340 345 350
 Tyr Val Phe Arg Asp Leu Ala Asn His Asn Gln Ile Phe Val Lys Asp
 355 360 365
 Lys Asp Pro Lys Val Tyr Asn Asn Pro Tyr Tyr Ile Asp Gln Val Gln
 370 375 380
 Leu Lys Asp Ala Gln Gln Ile Asp Leu Thr Ser Ile Gln Ala Ser Phe
 385 390 395 400
 Thr Thr Leu Asp Gly Val Asp Lys Thr Glu Ile Leu Lys Glu Leu Lys
 405 410 415
 Val Thr Asp Lys Asn Gln Asn Ala Ile Gln Ile Ser Asp Ile Thr Leu
 420 425 430
 Asp Thr Ser Lys Ser Leu Leu Ile Ile Lys Gly Asp Phe Asn Pro Lys
 435 440 445

Gln Gly His Phe Asn Ile Ser Tyr Asn Gly Asn Asn Val Met Thr Arg
 450 455 460
 Gln Ser Trp Glu Phe Lys Asp Gln Leu Tyr Ala Tyr Ser Gly Asn Leu
 465 470 475 480
 Gly Ala Val Leu Asn Gln Asp Gly Ser Lys Val Glu Ala Ser Leu Trp
 485 490 495
 Ser Pro Ser Ala Asp Ser Val Thr Met Ile Ile Tyr Asp Lys Asp Asn
 500 505 510
 Gln Asn Arg Val Val Ala Thr Thr Pro Leu Val Lys Asn Asn Lys Gly
 515 520 525
 Val Trp Gln Thr Ile Leu Asp Thr Lys Leu Gly Ile Lys Asn Tyr Thr
 530 535 540
 Gly Tyr Tyr Tyr Leu Tyr Glu Ile Lys Arg Gly Lys Asp Lys Val Lys
 545 550 555 560
 Ile Leu Asp Pro Tyr Ala Lys Ser Leu Ala Glu Trp Asp Ser Asn Thr
 565 570 575
 Val Asn Asp Asp Ile Lys Thr Ala Lys Ala Ala Phe Val Asn Pro Ser
 580 585 590
 Gln Leu Gly Pro Gln Asn Leu Ser Phe Ala Lys Ile Ala Asn Phe Lys
 595 600 605
 Gly Arg Gln Asp Ala Val Ile Tyr Glu Ala His Val Arg Asp Phe Thr
 610 615 620
 Ser Asp Arg Ser Leu Asp Gly Lys Leu Lys Asn Gln Phe Gly Thr Phe
 625 630 635 640
 Ala Ala Phe Ser Glu Lys Leu Asp Tyr Leu Gln Lys Leu Gly Val Thr
 645 650 655
 His Ile Gln Leu Pro Val Leu Ser Tyr Phe Tyr Val Asn Glu Met
 660 665 670
 Asp Lys Ser Arg Ser Thr Ala Tyr Thr Ser Ser Asp Asn Asn Tyr Asn
 675 680 685
 Trp Gly Tyr Asp Pro Gln Ser Tyr Phe Ala Leu Ser Gly Met Tyr Ser
 690 695 700
 Glu Lys Pro Lys Asp Pro Ser Ala Arg Ile Ala Glu Leu Lys Gln Leu
 705 710 715 720
 Ile His Asp Ile His Lys Arg Gly Met Gly Val Ile Leu Asp Val Val
 725 730 735
 Tyr Asn His Thr Ala Lys Thr Tyr Leu Phe Glu Asp Ile Glu Pro Asn
 740 745 750
 Tyr Tyr His Phe Met Asn Glu Asp Gly Ser Pro Arg Glu Ser Phe Gly
 755 760 765
 Gly Gly Arg Leu Gly Thr Thr His Ala Met Ser Arg Arg Val Leu Val
 770 775 780
 Asp Ser Ile Lys Tyr Leu Thr Ser Glu Phe Lys Val Asp Gly Phe Arg
 785 790 795 800
 Phe Asp Met Met Gly Asp His Asp Ala Ala Ala Ile Glu Leu Ala Tyr
 805 810 815
 Lys Glu Ala Lys Ala Ile Asn Pro Asn Met Ile Met Ile Gly Glu Gly
 820 825 830
 Trp Arg Thr Phe Gln Gly Asp Gln Gly Gln Pro Val Lys Pro Ala Asp
 835 840 845
 Gln Asp Trp Met Lys Ser Thr Asp Thr Val Gly Val Phe Ser Asp Asp
 850 855 860
 Ile Arg Asn Ser Leu Lys Ser Gly Phe Pro Asn Glu Gly Thr Pro Ala
 865 870 875 880
 Phe Ile Thr Gly Gly Pro Gln Ser Leu Gln Gly Ile Phe Lys Asn Ile
 885 890 895

Lys Ala Gln Pro Gly Asn Phe Glu Ala Asp Ser Pro Gly Asp Val Val
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 Lys Ser Ile
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<220>
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<400> 27

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cgt	gct	ggt	gat	act	gta	cgt	gtt	cac	gct	aaa	gtt	gaa	ggt	act	97	
Arg	Ala	Gly	Asp	Thr	Val	Arg	Val	His	Ala	Lys	Val	Val	Glu	Gly	Thr	
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cgc	gaa	cgt	att	cag	atc	ttt	gaa	ggt	gtt	atc	tca	cgt	aaa	ggt	145	
Arg	Glu	Arg	Ile	Gln	Ile	Phe	Glu	Gly	Val	Val	Ile	Ser	Arg	Lys	Gly	
35																45

caa	gga	atc	tca	gaa	atg	tac	aca	gta	cgt	aaa	att	tct	ggt	ggt	atc	193
Gln	Gly	Ile	Ser	Glu	Met	Tyr	Thr	Val	Arg	Lys	Ile	Ser	Gly	Gly	Ile	
50																60

ggt	gta	gag	cgt	aca	ttc	cca	att	cac	act	cct	cgt	gtt	gat	aaa	atc	241
Gly	Val	Glu	Arg	Thr	Phe	Pro	Ile	His	Thr	Pro	Arg	Val	Asp	Lys	Ile	
65																80

gaa	gtt	gtt	cgt	tat	ggt	aaa	gta	cgt	cgt	gct	aaa	ctt	tac	tac	tta	289
Glu	Val	Val	Arg	Tyr	Gly	Lys	Val	Arg	Arg	Ala	Lys	Leu	Tyr	Tyr	Leu	
85																95

cgc	gca	ttg	caa	ggtaaagctg	cacgttattaa	agaaatccgt	cgttaatttt	341
Arg	Ala	Leu	Gln					
100								

gatgatcaga	ttttaaaaat	gcttgggtgt	tttgggatag	taactatgtt	ttaaaactgg	401
acaaccaaga	cgtaaaaaat	ctgcctgtgg	gcagtttttt	tactaggatcc	ccttagttca	461
atggatataa	caactccctc	ctaaaggatg	attgtcggtt	cgattccggc	agggacata	521
ttcattgcat	gttaaatagcg	gttttagact	attttggccc	aaatttctct	gattaagttt	581
atcgttccta	tctttttgtt	cttgcatttg	atgtgcgtaa	acttctaaag	tgatattaa	641
attctcgtga	tctaaaactt	gagagatgga	aatttagatag	cttgcataatg	tatgcctgag	701
agagtgcact	cgtacctcgc	gaccagttt	ttttcgatata	gttttatgtt	ctgcattatt	761
tgaaagttt	tcgaataatc	tgtcggtttt	attttttgtt	aattcatgca	aaaaaaaataa	821
tgtatcattt	tcaatttggta	tattttctgtat	actactttttt	ttttttttttt	gcaggtatct	881
ttgggtgaaa	tgataatccc	aagttttattt	aattgataaa	tattttttttt	tgtaatcaat	941
atcatatcaat	gtttaaaccta	aacattcagc	gaagcgcattt	ccagttttttt	cgatgaggta	1001

aaaaagtcat	aaataaaa	ttataggaat	ctttctgg	aaaaaacagg	aaacatatac	4421
aggattatca	tctgatttta	gcgaaaat	ggttttgt	gattattcaa	ctagccaaga	4481
aatattaaat	aaatcagaga	ataatagaat	tgcaaaataa	attttaatgt	attctggtag	4541
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gtattctatt	aagaaagata	ataaaagcatt	cgaagagtct	ttagagtcag	tgagtggaa	4661
aaaacatata	attaaaataa	tgacttattc	gattatgtt	ggtggaaat	ttgttcttc	4721
attaatctt	g	ttagagaaag	aatttatgaa	ataggtat	ttttatctat	4781
tgaaacaact	aagatacaaa	ttataaggc	atttatattt	gagttat	tcatatcaat	4841
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attttaac	tcagagaact	caatgattt	cggtggaa	ttataaaata	aaagcagt	4961
tatgttaaac	ataacaacac	ttgcagaaag	ttat	ttataaagta	ttatgtttt	5021
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tgataaagtt	caaacagggaa	aaatctt	gtt	taagaatgaa	gatataaaaa	5321
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tttatcgccg	attgaaaata	ttagact	aaat	aaatca	gtagatgaga	5441
cgaatttagt	ttagataaaa	aacaataaa	aagaatgtt	atgaaattat	ctggtgg	5501
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<211> 111
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 <213> streptococcus

<400> 28

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						20				25			30		
Arg	Glu	Arg	Ile	Gln	Ile	Phe	Glu	Gly	Val	Val	Ile	Ser	Arg	Lys	Gly
						35				40			45		
Gln	Gly	Ile	Ser	Glu	Met	Tyr	Thr	Val	Arg	Lys	Ile	Ser	Gly	Gly	Ile
					50				55			60			
Gly	Val	Glu	Arg	Thr	Phe	Pro	Ile	His	Thr	Pro	Arg	Val	Asp	Lys	Ile
					65				70			75			80
Glu	Val	Val	Arg	Tyr	Gly	Lys	Val	Arg	Arg	Ala	Lys	Leu	Tyr	Tyr	Leu
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Arg	Ala	Leu	Gln	Gly	Lys	Ala	Ala	Arg	Ile	Lys	Glu	Ile	Arg	Arg	
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<210> 29

<211> 173
 <212> PRT
 <213> streptococcus

<400> 29

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						20				25			30		
Gln	Arg	Tyr	Leu	Pro	Thr	Lys	Asn	Lys	Ser	Ser	Ile	Arg	Asn	Ile	Pro
						35				40			45		
Ile	Asp	Asn	Asp	Thr	Leu	Phe	Phe	Leu	His	Glu	Phe	Thr	Lys	Asn	Lys

	50	55	60												
Asn	Asp	Arg	Leu	Phe	Asp	Lys	Leu	Ser	Asn	Asn	Ala	Val	Asn	Lys	Thr
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Ile	Arg	Lys	Ile	Thr	Gly	Arg	Glu	Val	Arg	Val	His	Ser	Leu	Arg	His
					85				90					95	
Thr	Phe	Ala	Ser	Tyr	Leu	Ile	Ser	Ile	Ser	Gln	Val	Leu	Asp	His	Glu
					100				105					110	
Asn	Leu	Asn	Ile	Thr	Leu	Glu	Val	Tyr	Ala	His	Gln	Leu	Gln	Glu	Gln
					115				120					125	
Lys	Asp	Arg	Asn	Asp	Lys	Leu	Asn	Gln	Arg	Asn	Leu	Gly	Gln	Asn	Ser
					130				135					140	
Ser	Lys	Pro	Leu	Phe	Thr	Cys	Asn	Glu	Tyr	Val	Pro	Cys	Arg	Asn	Arg
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Thr	Ser	Asn	Tyr	Ser	Leu	Gly	Gly	Ser	Cys	Tyr	Ile	His			
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<212> PRT
<213> streptococcus

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 35 40 45
 Gln Tyr Asp Gly Leu Ala Lys Leu Lys Asp Leu Lys Val Val Ser Gly
 50 55 60
 Glu Gln Ser Ile Asn Arg Glu Asp Leu Ser Asp Glu Phe Lys Asn Val
 65 70 75 80
 Val Ser Leu Glu Ala Thr Ser Asn Thr Lys Arg Asn Leu Leu Phe Ser
 85 90 95
 Ser Gly Val Phe Ser Phe Lys Glu Gly Lys Asn Ile Glu Glu Asn Asp
 100 105 110
 Lys Asn Ser Ile Leu Val His Glu Glu Phe Ala Lys Gln Asn Lys Leu
 115 120 125
 Lys Leu Gly Asp Glu Ile Asp Leu Glu Leu Leu Asp Thr Glu Lys Ser
 130 135 140
 Gly Lys Ile Lys Ser His Lys Phe Lys Ile Ile Gly Ile Phe Ser Gly
 145 150 155 160
 Lys Lys Gln Glu Thr Tyr Thr Gly Leu Ser Ser Asp Phe Ser Glu Asn
 165 170 175
 Met Val Phe Val Asp Tyr Ser Thr Ser Gln Glu Ile Leu Asn Lys Ser
 180 185 190
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 Ser Ile Met Leu Gly Gly Ile Val Val Leu Ser Leu Ile Leu Ile Leu
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Trp Leu Arg Glu Arg Ile Tyr Glu Ile Gly Ile Phe Leu Ser Ile Gly
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 Phe Gly Gly Ser Leu Ile Asn Lys Ser Ser Phe Met Leu Asn Ile Thr
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 Thr Leu Ala Glu Ser Tyr Leu Ile Leu Ile Ser Ile Ile Val Leu Ser
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 <211> 169
 <212> PRT
 <213> streptococcus

<400> 31
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 Lys Asn Glu Asp Ile Glu Lys Lys Gly Tyr Ser Asn His Arg Lys Asn
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<210> 32
 <211> 4171
 <212> DNA
 <213> Streptococcus

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 <211> 649
 <212> PRT
 <213> Streptococcus

<400> 33															
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					20				25				30		
His	Ala	Asp	Asn	His	Tyr	His	Phe	Phe	Asn	Gly	Lys	Ser	Leu	Ala	Thr
					35			40				45			
Phe	Asn	Thr	Asn	Gln	Leu	Ile	Arg	Glu	Val	Val	Tyr	Val	Glu	Ile	Ser
					50			55			60				
Leu	Asp	Thr	Met	Ser	Ser	Gly	Glu	His	Asp	Leu	Val	Lys	Val	Asn	Ile
					65			70			75			80	
Ile	Arg	Pro	Thr	Thr	Glu	His	Thr	Ile	Pro	Thr	Met	Met	Thr	Ala	Ser
					85				90			95			
Pro	Tyr	His	Gln	Gly	Ile	Asn	Asp	Pro	Ala	Ala	Asp	Gln	Lys	Thr	Tyr
					100			105			110				
Gln	Met	Glu	Gly	Ala	Leu	Ala	Val	Lys	Gln	Pro	Lys	His	Ile	Gln	Val
					115			120			125				
Asp	Thr	Lys	Pro	Phe	Lys	Glu	Glu	Val	Lys	His	Pro	Ser	Lys	Leu	Pro
					130			135			140				
Ile	Ser	Pro	Ala	Thr	Glu	Ser	Phe	Thr	His	Ile	Asp	Ser	Tyr	Ser	Leu
					145			150			155			160	
Asn	Asp	Tyr	Phe	Leu	Ser	Arg	Gly	Phe	Ala	Asn	Ile	Tyr	Val	Ser	Gly
					165				170			175			
Val	Gly	Thr	Ala	Gly	Ser	Thr	Gly	Phe	Met	Thr	Ser	Gly	Asp	Tyr	Gln
					180				185			190			
Gln	Ile	Gln	Ser	Phe	Lys	Ala	Val	Ile	Asp	Trp	Leu	Asn	Gly	Lys	Val
					195			200			205				
Thr	Ala	Phe	Thr	Ser	His	Lys	Arg	Asp	Lys	Gln	Val	Lys	Ala	Asp	Trp
					210			215			220				
Ser	Asn	Gly	Leu	Val	Ala	Thr	Thr	Gly	Lys	Ser	Tyr	Leu	Gly	Thr	Met
					225			230			235			240	
Ser	Thr	Gly	Leu	Ala	Thr	Thr	Gly	Val	Glu	Gly	Leu	Lys	Val	Ile	Ile
					245				250			255			
Ala	Glu	Ala	Ala	Ile	Ser	Thr	Trp	Tyr	Asp	Tyr	Tyr	Arg	Glu	Asn	Gly
					260			265			270				
Leu	Val	Cys	Ser	Pro	Gly	Gly	Tyr	Pro	Gly	Glu	Asp	Leu	Asp	Val	Leu
					275			280			285				
Thr	Glu	Leu	Thr	Tyr	Ser	Arg	Asn	Leu	Leu	Ala	Gly	Asp	Tyr	Ile	Lys
					290			295			300				
Asn	Asn	Asp	Cys	Tyr	Gln	Ala	Leu	Leu	Asn	Glu	Gln	Ser	Lys	Ala	Ile

305	310	315	320
Asp Arg Gln Ser Gly Asp Tyr Asn Gln Tyr Trp His Asp Arg Asn Tyr			
325	330	335	
Leu Thr His Val Asn Asn Val Lys Ser Arg Val Val Tyr Thr His Gly			
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Leu Gln Asp Trp Asn Val Lys Pro Arg His Val Tyr Lys Val Phe Asn			
355	360	365	
Ala Leu Pro Gln Thr Ile Lys Lys His Leu Phe Leu His Gln Gly Gln			
370	375	380	
His Val Tyr Met His Asn Trp Gln Ser Ile Asp Phe Arg Glu Ser Met			
385	390	395	400
Asn Ala Leu Leu Ser Gln Glu Leu Leu Gly Ile Asp Asn His Phe Gln			
405	410	415	
Leu Glu Glu Val Ile Trp Gln Asp Asn Thr Thr Glu Gln Thr Trp Gln			
420	425	430	
Val Leu Asp Ala Phe Gly Gly Asn His Gln Glu Gln Ile Gly Leu Gly			
435	440	445	
Asp Ser Lys Lys Leu Ile Asp Asn His Tyr Asp Lys Glu Ala Phe Asp			
450	455	460	
Thr Tyr Cys Lys Asp Phe Asn Val Phe Lys Asn Asp Leu Phe Lys Gly			
465	470	475	480
Asn Asn Lys Thr Asn Gln Ile Thr Ile Asn Leu Pro Leu Lys Lys Asn			
485	490	495	
Tyr Leu Leu Asn Gly Gln Cys Lys Leu His Leu Arg Val Lys Thr Ser			
500	505	510	
Asp Lys Ala Ile Leu Ser Ala Gln Ile Leu Asp Tyr Gly Pro Lys			
515	520	525	
Lys Arg Phe Lys Asp Thr Pro Thr Ile Lys Phe Leu Asn Ser Leu Asp			
530	535	540	
Asn Gly Lys Asn Phe Ala Arg Glu Ala Leu Arg Glu Leu Pro Phe Thr			
545	550	555	560
Lys Asp His Tyr Arg Val Ile Ser Lys Gly Val Leu Asn Leu Gln Asn			
565	570	575	
Arg Thr Asp Leu Leu Thr Ile Glu Ala Ile Glu Pro Glu Gln Trp Phe			
580	585	590	
Asp Ile Glu Phe Ser Leu Gln Pro Ser Ile Tyr Gln Leu Ser Lys Gly			
595	600	605	
Asp Asn Leu Arg Ile Ile Leu Tyr Thr Thr Asp Phe Glu His Thr Ile			
610	615	620	
Arg Asp Asn Ala Ser Tyr Ser Ile Thr Val Asp Leu Ser Gln Ser Tyr			
625	630	635	640
Leu Thr Ile Pro Thr Asn Gln Gly Asn			
645			

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 <212> PRT
 <213> Streptococcus

<400> 34			
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20	25	30	
His Asp Asn Lys Asn Ile Pro Gln Val Val Ala Thr Ile Val Asp Asp			
35	40	45	

56 / 63

Leu Gln Gly Ser Gly Ser Ser Asn His Phe Trp Tyr Phe Gly Asn Thr
 50 55 60
 Thr Asp Thr Ser Ile Leu Met Ile Ala His Leu Asn Arg Lys Phe Tyr
 65 70 75 80
 Ile Gln Val Asn Leu Lys Asp Phe Asp Phe Ala Leu Asn Leu Ile Ala
 85 90 95
 Ile Asn Asn Trp Lys Ser Leu Leu Gln Thr Gln Leu Glu Ala Leu Asn
 100 105 110
 Asp Thr Leu Ala Ile Phe Gln
 115

<210> 35
 <211> 326
 <212> PRT
 <213> Streptococcus

<400> 35
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 Ile Glu Ala Ala Leu Ser Gln Leu Thr Ala Ala Gly Gly Lys Gln Leu
 35 40 45
 Arg Pro Ala Phe Phe Tyr Leu Phe Ser Gln Leu Gly Asn Lys Glu Asn
 50 55 60
 Gln Asp Thr Gln Gln Leu Lys Lys Ile Ala Ala Ser Leu Glu Ile Leu
 65 70 75 80
 His Val Ala Thr Leu Ile His Asp Asp Val Ile Asp Asp Ser Pro Leu
 85 90 95
 Arg Arg Gly Asn Met Thr Ile Gln Ser Lys Phe Gly Lys Asp Ile Ala
 100 105 110
 Val Tyr Thr Gly Asp Leu Leu Phe Thr Val Phe Phe Asp Leu Ile Leu
 115 120 125
 Glu Ser Met Thr Asp Thr Pro Phe Met Arg Ile Asn Ala Lys Ser Met
 130 135 140
 Arg Lys Ile Leu Met Gly Glu Leu Asp Gln Met His Leu Arg Tyr Asn
 145 150 155 160
 Gln Gln Gln Gly Ile His His Tyr Leu Arg Ala Ile Ser Gly Lys Thr
 165 170 175
 Ala Glu Leu Phe Lys Leu Ala Ser Lys Glu Gly Ala Tyr Phe Gly Gly
 180 185 190
 Ala Glu Lys Glu Val Val Arg Leu Ala Gly His Ile Gly Phe Asn Ile
 195 200 205
 Gly Met Thr Phe Gln Ile Leu Asp Asp Ile Leu Asp Tyr Thr Ala Asp
 210 215 220
 Lys Lys Thr Phe Asn Lys Pro Val Leu Glu Asp Leu Thr Gln Gly Val
 225 230 235 240
 Tyr Ser Leu Pro Leu Leu Ala Ile Glu Glu Asn Pro Asp Ile Phe
 245 250 255
 Lys Pro Ile Leu Asp Lys Lys Thr Asp Met Ala Thr Glu Asp Met Glu
 260 265 270
 Lys Ile Ala Tyr Leu Val Val Ser His Arg Gly Val Asp Lys Ala Arg
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 His Leu Ala Arg Lys Phe Thr Glu Lys Ala Ile Ser Asp Ile Asn Lys
 290 295 300
 Leu Pro Gln Asn Ser Ala Lys Lys Gln Leu Leu Gln Leu Thr Asn Tyr

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Leu	Leu
Lys	Arg
	Lys
	Ile
	325

315	320
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<210> 36
 <211> 247
 <212> PRT
 <213> Streptococcus

<400> 36

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 Lys Glu Gly Glu Lys Ile Ala Ile Leu Gly Arg Ser Gly Ser Gly Lys
 35 40 45
 Ser Thr Leu Ala Ser Leu Leu Arg Gly Asp Leu Lys Ala Ser Gln Gly
 50 55 60
 Lys Ile Thr Leu Gly Gly Ala Asp Val Ser Ile Val Gly Asp Cys Ile
 65 70 75 80
 Ser Asn Tyr Ile Gly Val Ile Gln Gln Ala Pro Tyr Leu Phe Asn Thr
 85 90 95
 Thr Leu Leu Asn Asn Ile Arg Ile Gly Asn Gln Asp Ala Ser Glu Glu
 100 105 110
 Asp Val Trp Lys Val Leu Glu Arg Val Gly Leu Lys Glu Met Val Thr
 115 120 125
 Asp Leu Ser Asp Gly Leu Tyr Thr Met Val Asp Glu Ala Gly Leu Arg
 130 135 140
 Phe Ser Gly Gly Glu Arg His Arg Ile Ala Leu Ala Arg Ile Leu Leu
 145 150 155 160
 Lys Asp Val Pro Ile Val Ile Leu Asp Glu Pro Thr Val Gly Leu Asp
 165 170 175
 Pro Ile Thr Glu Gln Ala Leu Leu Arg Val Phe Met Lys Glu Leu Glu
 180 185 190
 Gly Lys Thr Leu Val Trp Ile Thr His His Leu Lys Gly Ile Glu His
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 Ala Asp Arg Ile Leu Phe Ile Glu Asn Gly Gln Leu Glu Leu Glu Gly
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 Ser Pro Gln Glu Leu Ser Gln Ser Ser Gln Arg Tyr Arg Gln Leu Lys
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 Ala Ala Asp Asp Gly Asp Leu
 245

<210> 37
 <211> 3480
 <212> DNA
 <213> Streptococcus

<400> 37

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caaatacatgt	ttcatatgtt	atctggcaac	aaaagtttta	ctcaaaataca	aatatgtattt	2280
atggaccctgc	taataacttgg	aatgaatgc	cagatcgtgg	ttggcgttact	gccaaaccatt	2340
atgaccatgt	tcacgtatca	tttacaaaat	aatataaaaa	aggaagctat	ttggcgttctt	2400
tttttatatgc	tttgaataga	ctttaaagg	tcttatctaa	tttttattaa	attgaggaga	2460
ttaagctata	agtctgaaac	tactttcagc	ttaaccgtga	ctaaatcaaa	acgttaaaaac	2520
taaaaatctaa	gtctgtaaag	attattgaaa	acgtttaaa	aacagatata	ataaggttt	2580
tagatatcta	aaattaaaaaa	agataaggaa	gtgagaatat	gccacatcta	agtaaaaagaag	2640
cttttaaaaaa	gcaaaaaaaa	aatggcatta	tttggtgtat	tcaagcttt	cctggggagc	2700
ctctttatac	tgaaagtgg	ggtggatgtc	ctcttttagc	tttggcagct	caagaagcag	2760
gagcggttgg	tataagagcc	aatagtgtcc	gcccacattaa	ggaaattcaa	gaagttacta	2820
atttacctat	catcgccatt	attaaacgt	aatatcc	acaagaacca	tttacactg	2880
ctacgatgac	agaggtggat	caattagct	gtttagat	tgccatata	gccttagatt	2940
gtacacttag	agagcgctat	gatgggttga	gtgttagct	gtttattca	agataaaaag	3000
ggaaaatatcc	tgaacagtt	ctaattggct	atataagtac	tttgaagaa	ggtaaaaatg	3060
cttttgaagc	aggagttgtat	tttggggta	caactctatc	tggatcacaca	gattacagcc	3120
gccaagaaga	aggacccggat	atagaactcc	ttaataagct	ttgtcaagcc	ggtatagatg	3180
tgattgcgg	aggtaaaatt	catactcc	agcaagctaa	tgaattaaat	catataaggtg	3240
ttgcaggaat	tgttagttgt	ggtgcgtatca	ctagacaaa	agaaatagcg	gagcgttca	3300
tctcaggact	tagttaaaag	tgttactcaa	aaatcaaaaat	caaaaataaaa	aagggaata	3360
gttatgagta	tcaaaaaaaag	tgtgattgg	tttgcctcg	gagctgcagc	attatcaatg	3420
tttgcttgcgt	tagacagtag	tcaatctgtt	atggcgtccg	agaaggataa	agtcgaaatt	3480

<210> 38
 <211> 306
 <212> PRT
 <213> Streptococcus

<400> 38

Asn	Ser	Ile	Trp	Arg	Phe	Phe	Leu	Asn	Lys	Trp	Leu	Val	Lys	Ala	Ser
1				5					10						15
Ser	Leu	Val	Val	Leu	Gly	Gly	Met	Val	Leu	Ser	Ala	Gly	Ser	Arg	Val
					20			25						30	
Leu	Ala	Asp	Thr	Tyr	Val	Arg	Pro	Ile	Asp	Asn	Gly	Arg	Ile	Thr	Thr
					35			40				45			
Gly	Phe	Asn	Gly	Tyr	Pro	Gly	His	Cys	Gly	Val	Asp	Tyr	Ala	Val	Pro
					50			55			60				
Thr	Gly	Thr	Ile	Ile	Arg	Ala	Val	Ala	Asp	Gly	Thr	Val	Lys	Phe	Ala
					65			70			75			80	
Gly	Ala	Gly	Ala	Asn	Phe	Ser	Trp	Met	Thr	Asp	Leu	Ala	Gly	Asn	Cys
					85				90			95			
Val	Met	Ile	Gln	His	Ala	Asp	Gly	Met	His	Ser	Gly	Tyr	Ala	His	Met
					100				105			110			
Ser	Arg	Val	Val	Ala	Arg	Thr	Gly	Glu	Lys	Val	Lys	Gln	Gly	Asp	Ile
					115			120			125				
Ile	Gly	Tyr	Val	Gly	Ala	Thr	Gly	Met	Ala	Thr	Gly	Pro	His	Leu	His
					130			135			140				
Phe	Glu	Phe	Leu	Pro	Ala	Asn	Pro	Asn	Phe	Gln	Asn	Gly	Phe	His	Gly
					145			150			155			160	
Arg	Ile	Asn	Pro	Thr	Ser	Leu	Ile	Ala	Asn	Val	Ala	Thr	Phe	Ser	Gly
					165				170			175			
Lys	Thr	Gln	Ala	Ser	Ala	Pro	Ser	Ile	Lys	Pro	Leu	Gln	Ser	Ala	Pro
					180				185			190			
Val	Gln	Asn	Gln	Ser	Ser	Lys	Leu	Lys	Val	Tyr	Arg	Val	Asp	Glu	Leu
					195			200			205				
Gln	Lys	Val	Asn	Gly	Val	Trp	Leu	Val	Lys	Asn	Asn	Thr	Leu	Thr	Pro
					210			215			220				
Thr	Gly	Phe	Asp	Trp	Asn	Asp	Asn	Gly	Ile	Pro	Ala	Ser	Glu	Ile	Asp
					225			230			235			240	
Glu	Val	Asp	Ala	Asn	Gly	Asn	Leu	Thr	Ala	Asp	Gln	Val	Leu	Gln	Lys
					245				250			255			
Gly	Gly	Tyr	Phe	Ile	Phe	Asn	Pro	Lys	Thr	Leu	Lys	Thr	Val	Glu	Lys
					260			265			270				
Pro	Ile	Gln	Gly	Thr	Ala	Gly	Leu	Thr	Trp	Ala	Lys	Thr	Arg	Phe	Ala
					275			280			285				
Asn	Gly	Ser	Ser	Val	Trp	Leu	Arg	Val	Asp	Asn	Ser	Gln	Glu	Leu	Leu
					290			295			300				
Tyr	Lys														
		305													

<210> 39
 <211> 434
 <212> PRT
 <213> Streptococcus

<400> 39

Met	Lys	Met	Asn	Lys	Lys	Val	Leu	Leu	Thr	Ser	Thr	Met	Ala	Ala	Ser
1				5				10						15	
Leu	Leu	Ser	Val	Ala	Ser	Val	Gln	Ala	Gln	Glu	Thr	Asp	Thr	Thr	Trp
					20			25			30				
Thr	Ala	Arg	Thr	Val	Ser	Glu	Val	Lys	Ala	Asp	Leu	Val	Lys	Gln	Asp
					35			40			45				
Asn	Lys	Ser	Ser	Tyr	Thr	Val	Lys	Tyr	Gly	Asp	Thr	Leu	Ser	Val	Ile

50	55	60
Ser Glu Ala Met Ser Ile Asp Met Asn Val Leu Ala Lys Ile Asn Asn		
65	70	75
Ile Ala Asp Ile Asn Leu Ile Tyr Pro Glu Thr Thr Leu Thr Val Thr		80
85	90	95
Tyr Asp Gln Lys Ser His Thr Ala Thr Ser Met Lys Ile Glu Thr Pro		
100	105	110
Ala Thr Asn Ala Ala Gly Gln Thr Thr Ala Thr Val Asp Leu Lys Thr		
115	120	125
Asn Gln Val Ser Val Ala Asp Gln Lys Val Ser Leu Asn Thr Ile Ser		
130	135	140
Glu Gly Met Thr Pro Glu Ala Ala Thr Thr Ile Val Ser Pro Met Lys		
145	150	155
Thr Tyr Ser Ser Ala Pro Ala Leu Lys Ser Lys Glu Val Leu Ala Gln		
165	170	175
Glu Gln Ala Val Ser Gln Ala Ala Asn Glu Gln Val Ser Thr Ala		
180	185	190
Pro Val Lys Ser Ile Thr Ser Glu Val Pro Ala Ala Lys Glu Glu Val		
195	200	205
Lys Pro Thr Gln Thr Ser Val Ser Gln Ser Thr Thr Val Ser Pro Ala		
210	215	220
Ser Val Ala Ala Glu Thr Pro Ala Pro Val Ala Lys Val Ala Pro Val		
225	230	235
Arg Thr Val Ala Ala Pro Arg Val Ala Ser Val Lys Val Val Thr Pro		
245	250	255
Lys Val Glu Thr Gly Ala Ser Pro Glu His Val Ser Ala Pro Ala Val		
260	265	270
Pro Val Thr Thr Ser Thr Ala Thr Asp Ser Lys Leu Gln Ala Thr		
275	280	285
Glu Val Lys Ser Val Pro Val Ala Gln Lys Ala Pro Thr Ala Thr Pro		
290	295	300
Val Ala Gln Pro Ala Ser Thr Thr Asn Ala Val Ala Ala His Pro Glu		
305	310	315
Asn Ala Gly Leu Gln Pro His Val Ala Ala Tyr Lys Glu Lys Val Ala		
325	330	335
Ser Thr Tyr Gly Val Asn Glu Phe Ser Thr Tyr Arg Ala Gly Asp Pro		
340	345	350
Gly Asp His Gly Lys Gly Leu Ala Val Asp Phe Ile Val Gly Lys Asn		
355	360	365
Gln Ala Leu Gly Asn Glu Val Ala Gln Tyr Ser Thr Gln Asn Met Ala		
370	375	380
Ala Asn Asn Ile Ser Tyr Val Ile Trp Gln Gln Lys Phe Tyr Ser Asn		
385	390	395
Thr Asn Ser Ile Tyr Gly Pro Ala Asn Thr Trp Asn Ala Met Pro Asp		
405	410	415
Arg Gly Gly Val Thr Ala Asn His Tyr Asp His Val His Val Ser Phe		
420	425	430
Asn Lys		

<210> 40
 <211> 232
 <212> PRT
 <213> Streptococcus

<400> 40

Met Pro His Leu Ser Lys Glu Ala Phe Lys Lys Gln Ile Lys Asn Gly
 1 5 10 15
 Ile Ile Val Ser Cys Gln Ala Leu Pro Gly Glu Pro Leu Tyr Thr Glu
 20 25 30
 Ser Gly Gly Val Met Pro Leu Leu Ala Ala Gln Glu Ala Gly
 35 40 45
 Ala Val Gly Ile Arg Ala Asn Ser Val Arg Asp Ile Lys Glu Ile Gln
 50 55 60
 Glu Val Thr Asn Leu Pro Ile Ile Gly Ile Ile Lys Arg Glu Tyr Pro
 65 70 75 80
 Pro Gln Glu Pro Phe Ile Thr Ala Thr Met Thr Glu Val Asp Gln Leu
 85 90 95
 Ala Ser Leu Asp Ile Ala Val Ile Ala Leu Asp Cys Thr Leu Arg Glu
 100 105 110
 Arg His Asp Gly Leu Ser Val Ala Glu Phe Ile Gln Lys Ile Lys Gly
 115 120 125
 Lys Tyr Pro Glu Gln Leu Leu Met Ala Asp Ile Ser Thr Phe Glu Glu
 130 135 140
 Gly Lys Asn Ala Phe Glu Ala Gly Val Asp Phe Val Gly Thr Thr Leu
 145 150 155 160
 Ser Gly Tyr Thr Asp Tyr Xaa Arg Gln Glu Glu Gly Pro Asp Ile Glu
 165 170 175
 Leu Leu Asn Lys Leu Cys Gln Ala Gly Ile Asp Val Ile Ala Glu Gly
 180 185 190
 Lys Ile His Thr Pro Lys Gln Ala Asn Glu Ile Asn His Ile Gly Val
 195 200 205
 Ala Gly Ile Val Val Gly Gly Ala Ile Thr Arg Pro Lys Glu Ile Ala
 210 215 220
 Glu Arg Phe Ile Ser Gly Leu Ser
 225 230

<210> 41
 <211> 39
 <212> PRT
 <213> Streptococcus

<400> 41
 Met Ser Ile Lys Lys Ser Val Ile Gly Phe Cys Leu Gly Ala Ala Ala
 1 5 10 15
 Leu Ser Met Phe Ala Cys Val Asp Ser Ser Gln Ser Val Met Ala Ala
 20 25 30
 Glu Lys Asp Lys Val Glu Ile
 35

<210> 42
 <211> 1305
 <212> DNA
 <213> Streptococcus

<400> 42
 atgaaaatga ataaaaaggt actattgaca tcgacaatgg cagttcgct attatcagtc 60
 gcaagtgttc aagcacaaga aacagatacg acgtggacag cacgtactgt ttcagaggtta 120
 aaggctgatt tggtaaagca agacaataaa tcatcatata ctgtgaaata tggtgataca 180
 ctaagcgtta tttcagaagc aatgtcaatt gatatgaatg tcttagcaaa aattaataac 240
 atgcagata tcaatcttat ttatcctgag acaacactga cagtaactta cgatcagaag 300
 agtcatactg ccacttcaat gaaaatagaa acaccagcaa caaatgctgc tggtcaaaca 360

acagctactg	tggatttcaa	aaccaatcaa	gtttctgttgc	cagacaaaaa	agtttctctc	420
aatacaattt	cggaaggat	gacaccagaa	gcagcaacaa	cgattgttc	gccaatgaag	480
acatattctt	ctgcgccagc	tttgaatca	aaagaagtt	tagcacaaga	gcaagctgtt	540
agtcaagcag	cagctaatga	acaggtatca	acagctcctg	tgaagtctat	tacttcagaa	600
gttccagcag	ctaaagagga	agttaaacca	actcagacgt	cagtcagtc	gtcaacaaca	660
gtatcaccag	cttctgttgc	cgctgaaaca	ccagctccag	tagctaaatg	agcaccggta	720
agaactgtag	cagccccctag	agtggcaagt	gtttaagtag	tcactcccaa	agtagaaact	780
ggtgcatacc	cagagcatgt	atcagtcctca	gcagttccctg	tgactacgc	ttcaacagct	840
acagacagta	agttacaagc	gactgaagtt	aagagcgttc	cggttagcaca	aaaagctcca	900
acagcaacac	cggtagcaca	accagttca	acaacaaatg	cagtagctgc	acatccctgaa	960
aatgcagggc	tccaaacctca	tgttgcagct	tataaagaaaa	aagtagcgtc	aacttatggaa	1020
gttaatgaat	tcagtagata	ccgtgcaggt	gatccaggtg	atcatggtaa	aggtttagca	1080
gtcgacttta	ttttaggtaa	aaaccaagca	cttggtaatg	aagttgcaca	gtactctaca	1140
caaaatatgg	cagcaaataa	catttcata	gttatctggc	aacaaaatgt	ttactcaaata	1200
acaaaatagta	tttatggacc	tgctaatact	tggaatgca	tgccagatcg	tggggcggtt	1260
actgccaacc	attatgacca	tgttacgt	tcatttaaca	aataaa		1305

<210> 43
 <211> 1230
 <212> DNA
 <213> Streptococcus

<400> 43						
caagaaacag	atacgacgt	gacagcacgt	actgtttcag	aggtaaaggc	tgatttggt	60
aagaagaca	ataaaatcatc	ataatactgt	aaatatggt	atacactaa	cgttatccaa	120
gaagcaatgt	caattgtat	gaatgttca	gcaaaaatta	ataacattgc	agatatcaat	180
cttatttac	ctgagacaac	actgacagta	attacatgc	agaagagtc	tactgcccact	240
tcaatgaaaa	tagaaaacacc	agcaacaaat	gtctgtggtc	aaacaacacg	tactgtggat	300
ttgaaaacca	atcaagttc	tgttgcagac	caaaaagttt	ctctcaatac	aatttcggaa	360
ggtatgacac	cagaaggcgc	aacaacgatt	gttgcgc	tgaagacata	ttcttctgcg	420
ccagcttga	aatcaaaaga	agtattagca	caagagcag	ctgttagtca	agcagcagct	480
aatgaacagg	tatcaacacg	tcctgtgaag	tcgattactt	cagaagtcc	agcagctaaa	540
gaggaagtt	aaccaactca	gacgtcagtc	agtcagtc	caacagtatc	accagcttct	600
gttgcgcgt	aaacaccagc	tccagtagct	aaagtagcac	cggtaaagac	tgtagcagcc	660
cctagagtgg	caagtgtt	aaagtagtact	cctaaatgt	aaactgtgc	atcaccagag	720
catgtatcag	ctccagcagt	tcctgtgact	acgacttca	cagctacaga	cagtaagtt	780
caagcactg	aagttaaagag	cgttccggta	gcacaaaaag	ctccaacagc	aacaccggta	840
gcacaaccag	cttcaacaaac	aaatgcagta	gtgcacatc	ctgaaaatgc	agggtccaa	900
cctcatgtt	cagcttataa	agaaaaatgt	gcgtcaactt	atggagttaa	tgaattcagt	960
acataccgtg	caggtgatcc	aggtgatcat	ggtaaagggtt	tagcagtc	ctttattgt	1020
ggtaaaaacc	aagcacttgg	taatgaagtt	gcacagtt	ctacacaaaa	tatggcagca	1080
aataacattt	cataatgtt	ctggcaacaa	aagttttact	caaataaaa	tagtatttt	1140
ggacctgtca	atacttggaa	tgcaatgca	gatcgtggt	gcgttactgc	caaccattat	1200
gaccatgttc	acgtatcatt	taacaaataa				1230

<210> 44
 <211> 409
 <212> PRT
 <213> Streptococcus

<400> 44															
Gln	Glu	Thr	Asp	Thr	Thr	Trp	Thr	Ala	Arg	Thr	Val	Ser	Glu	Val	Lys
1				5				10				15			
Ala	Asp	Leu	Val	Lys	Gln	Asp	Asn	Lys	Ser	Ser	Tyr	Thr	Val	Lys	Tyr
				20				25				30			
Gly	Asp	Thr	Leu	Ser	Val	Ile	Ser	Glu	Ala	Met	Ser	Ile	Asp	Met	Asn

35	40	45
Val Leu Ala Lys Ile Asn Asn Ile Ala Asp Ile Asn Leu Ile Tyr Pro		
50	55	60
Glu Thr Thr Leu Thr Val Thr Tyr Asp Gln Lys Ser His Thr Ala Thr		
65	70	75
Ser Met Lys Ile Glu Thr Pro Ala Thr Asn Ala Ala Gly Gln Thr Thr		
85	90	95
Ala Thr Val Asp Leu Lys Thr Asn Gln Val Ser Val Ala Asp Gln Lys		
100	105	110
Val Ser Leu Asn Thr Ile Ser Glu Gly Met Thr Pro Glu Ala Ala Thr		
115	120	125
Thr Ile Val Ser Pro Met Lys Thr Tyr Ser Ser Ala Pro Ala Leu Lys		
130	135	140
Ser Lys Glu Val Leu Ala Gln Glu Gln Ala Val Ser Gln Ala Ala Ala		
145	150	155
Asn Glu Gln Val Ser Thr Ala Pro Val Lys Ser Ile Thr Ser Glu Val		
165	170	175
Pro Ala Ala Lys Glu Glu Val Lys Pro Thr Gln Thr Ser Val Ser Gln		
180	185	190
Ser Thr Thr Val Ser Pro Ala Ser Val Ala Ala Glu Thr Pro Ala Pro		
195	200	205
Val Ala Lys Val Ala Pro Val Arg Thr Val Ala Ala Pro Arg Val Ala		
210	215	220
Ser Val Lys Val Val Thr Pro Lys Val Glu Thr Gly Ala Ser Pro Glu		
225	230	235
His Val Ser Ala Pro Ala Val Pro Val Thr Thr Thr Ser Thr Ala Thr		
245	250	255
Asp Ser Lys Leu Gln Ala Thr Glu Val Lys Ser Val Pro Val Ala Gln		
260	265	270
Lys Ala Pro Thr Ala Thr Pro Val Ala Gln Pro Ala Ser Thr Thr Asn		
275	280	285
Ala Val Ala Ala His Pro Glu Asn Ala Gly Leu Gln Pro His Val Ala		
290	295	300
Ala Tyr Lys Glu Lys Val Ala Ser Thr Tyr Gly Val Asn Glu Phe Ser		
305	310	315
Thr Tyr Arg Ala Gly Asp Pro Gly Asp His Gly Lys Gly Leu Ala Val		
325	330	335
Asp Phe Ile Val Gly Lys Asn Gln Ala Leu Gly Asn Glu Val Ala Gln		
340	345	350
Tyr Ser Thr Gln Asn Met Ala Ala Asn Asn Ile Ser Tyr Val Ile Trp		
355	360	365
Gln Gln Lys Phe Tyr Ser Asn Thr Asn Ser Ile Tyr Gly Pro Ala Asn		
370	375	380
Thr Trp Asn Ala Met Pro Asp Arg Gly Gly Val Thr Ala Asn His Tyr		
385	390	395
Asp His Val His Val Ser Phe Asn Lys		
405		